AEROSIL (R) FOR THE IMPROVEMENT OF THE FLOW BEHAVIOR OF POWDERED SUBSTANCES

Translation of "AEROSIL (R) zur Verbesserung des Fließverhaltens Pulverformiger Substanzen Degussa", Frankfurt, West Germany, No. 31, August 10, 1971, pp. 1-17.
This article is a study to investigate the problems related to the fact that powdered substances tend to agglomerate and stick together whatever their composition and use. This impedes the packing, storage and handling of these materials.

The study investigated AEROSIL (R) as an agent to improve the "free flowing" characteristics of powdered materials. A summary statement concludes, that the use of AEROSIL 200, AEROSIL R 972, aluminum oxide C and sylicic acid D 17 as flow agents has made broad improvements possible in the properties of powdered products.
The highly dispersed silicic acid AEROSIL (R) has been used for many applications in numerous branches of industry. In the course of developing new methods of producing silicic acids, new products have been introduced with unusual properties, especially for application in powdered materials.

Powdered substances tend to agglomerate and stick together whatever their composition and use. This impedes the packing, storage and handling of this material. However, increasing simplification in industry demands good flow behavior of these powdered materials as well as stability in storage, weathering and under pressure.

1. Survey of the literature

The attempts to improve the flow properties of powdered products are mirrored in a multiplicity of patents and treatises. However, these are mostly limited to the description of individual cases.

In this way the following products are named, for example, as "flow facilitators" or "flow helpers" for ammonium fertilizers. alipathic amine in combination with carrier materials hydrazine and guanidine compounds urea-formaldehyde resin solutions phthalocyanine compounds sprays of high molecular weight glycols paraffin sulfo-phenolic acid and alkyl-aryl-sulfonate fatty acid aluminum salts

*Numbers in margin indicate pagination of foreign text
chromium, cadmium and tin salts, also such simple organic acids including iron ammonium citrate
alkali- and alkaline earth phosphate
magnesium oxide and calcinated dolomite
hexacyanoferrate
copper-I-oxide, copper-II-oxide, also zirconium-oxychloride
boron trioxide
rice flour

This enumeration—only a selection from a far greater number of publications—shows the multiplicity of the recommended flow agents. These have been applied in different forms, such as: mixtures, sprays and coatings. Soon silicic acids and silicates found a more uniform and, therefore, more clearly arranged application. Of these, the type obtained from the gas phase, e.g., the AEROSIL types have especially found increasing use by manufacturers and processors of powdered products. Some examples from the literature are:
prevention of clumping of fertilizers (1)
improved flow of pickling salts (2)
flow behavior of foods, drugs and cosmetics (3)
grinding and free-flowing aid for butyl-hydroxyl-toluol, methionine and its calcium salts, sodium propionate, urea, vitamins, diet products, spices, etc. (4)
flow agent for sodium chlorate and sodium chloride (5)
for the loosening of malt and molasses preparations (6)
for improving the flow of powdered waves (e.g., Carbowachs 600) (7)

2. Basics of flow behavior of powders and the mode of action of flow agents

The flow behavior of powders is determined principally by the physical properties of the powder:
hygroscopicity or sensitivity to weathering
form and size distribution of the powder particles
ability to become electrostatically charged
sintering at low temperatures
adhesion of the particles to each other

Investigations of the influence of these factors and
suggestions for improving of flow behavior are displaced in
the newer literature in the following examples:

2.1. M. Frey (8) indicates that for crystalline products
even traces of moisture produce an extremely thin film of fluid,
often in the form of a saturated solution of the product.
This acts like a binding agent in causing the particles to
adhere and thus hinders their "free flowing".

2.2. R. Tawashi (9) reports about the influence of fric-
tion between the particles as a quantity obstructing the flow
behavior. He also mentions the possibility of forming a
"single particle" covering of the powder particles by means
of the addition of small amounts of highly dispersed flow
agents, e.g., sylicic acids. Such a covering of the powder
particles by a single layer hull of flow agent would allow a
minimal value to be achieved for the friction between differ-
ent particles. The optimal amount of flow agent can be cal-
culated as a function of the density and average particle size
(10).

2.3. F. Gstirner and C. Pick (11) refer to the differ-
ence between specific heavy powdered material with good flow
properties and specific light powder materials which mostly
flow poorly. For the latter the ability to flow can be im-
proved by means of highly dispersed flow agents from 2 to 8
times.

2.4. W. A. Ritschel (12) describes the electrostatic
charging of powders as a cause of disturbance in their flow
behavior and of the appearance of reagglomeration on grinding
of such materials. He specifies the addition of highly dis-
persed sylicic acid as a remedy for this.

2.5. H. von Czetsch-Lindenwald and A. F. Asker (13) reports about the use of flow agents in improving the measure-
ment of dosage in medicinal materials for the correct filling
of capsules. Here, too, sylicic acids are recommended as an
effective flow agent.

3. AEROSIL as a flow agent

The following demands for an effective flow agent with
the most general possible application result from investigations
of the theory of flow behavior.

3.1 Extreme particle fineness for the single layer cover-
ing of the material used with only small additions. Figure 1
shows the particle fineness of AEROSIL 200 by means of an
electron micrograph.

3.2 The loosest possible agglomeration of the primary
particles which largely also divides into the actual effective
primary particle form or minimal agglomeration through simple
mixing procedures.

3.3 Largely inactive chemically with the materials to be
used.

3.4 High chemical purity and no physiological problems
for application for pharmaceuticals, human and animal food.

3.5. Absolute colorlessness, so that no discoloration of
the substances to be used need be feared.
3.6 Hydrophilic or hydrophobic surface properties choosable for specific cases.

The demands named under 3.1 and 3.6 have been fulfilled by the highly dispersed sylicic acids and aluminum oxide produced according to the AEROSIL procedures (14). In the past years AEROSIL has increasingly been used as a grinding and flow agent. It is produced from silicon tetrachloride by flame hydrolysis and is the purest form of sylicic acid produced large scale (15).

In addition to a high degree of chemical purity, hygienic harmlessness, high loose volume and extreme particle fineness are outstanding properties of AEROSIL for its use as a flow agent. The particle size distribution of AEROSIL 200 is shown in Figure 2. As a consequence of its high absorbptive capacity for fluids of all types, AEROSIL is appropriate for the conversion of liquid, plastic lubricating or agglomerated mixtures of materials to "dry" largely pressure stable and free flowing powdered products. Of the different kinds of AEROSIL, AEROSIL 200 is recommended for this application. AEROSIL (R) 972 is a special type AEROSIL and differs principally in its surface properties. While AEROSIL 200, like all the usual sylicic acids, is hydrophilic, AEROSIL R 972 is hydrophobic. This is brought about by replacing the sinanol groups on its surface with di-methyl-dichlorosilane (16).

A consequence of the hydrophobic character of AEROSIL R 972 is its ombrophobicness in water. In many cases, this is of decisive significance for its outstanding effect as a flow agent in comparison with ordinary hydrophilic sylicic acids. Its use as a flow agent represent considerable progress for the methods of improving the flow behavior of powdered materials.
Analogously to the AEROSIL procedure, aluminum oxide C is also produced from aluminum chloride by flame hydrolysis and, according to current experience, is principally useful in reducing the tendency of powdered materials to acquire electrostatic charge.

4. **Mixing technique**

The special properties of AEROSIL 200 and AEROSIL R 972 as flow agents (especially AEROSIL R 972), namely, extreme particle fineness and loose secondary structure, can also be optimally effective if the method of mixing leads to a good coating of the powdered particles by the AEROSIL.

According to extensive investigations, the following method of mixing is the most effective:

All the AEROSIL is placed in a mixer which operates according to the free fall principle, and the powdered product to be coated is added in approximately 4 or 5 portions under continuous mixing. The distribution of the AEROSIL then succeeds according to the model of the origin of a "snowball" and can be determined in the following example both mathematically and experimentally.

For an addition of 1% AEROSIL with an average effective secondary particle size of 0.5 microns (taken as an example) to sodium hydrogen carbonate, more than 1000 secondary particles of AEROSIL are available for coating each sodium hydrogen carbonate particle. In this only the experimentally useful secondary structure of the AEROSIL is employed and not its primary particle sizes since here more than 10 million AEROSIL
primary particles would be available for coating each sodium hydrogen carbonate particle.

The degree of agreement between this mathematical example and practice is shown in the following experiment in which hydrophobic AEROSIL R 972 is used in coating the particles for better visibility:

Procedure: while the untreated sodium hydrogen carbonate is quickly wet by water and sinks, the product which was treated with 1% AEROSIL R 972 remains persistently on the surface of the water without becoming wet (see Figure 3).

This method of mixing according to the "snowball principle" is based on the following processes:

With the addition of the first fifth of the powder being treated into the entire amount of the AEROSIL applied, "snowballs" form a loose, at first still surplus sheet of "snow" (AEROSIL) around the unequally large particles of the powder. A surplus of AEROSIL also remains. With the addition of the next fifth, the newly added powder particles take the AEROSIL necessary for coating them from the still available surplus AEROSIL. In the final state of the mixture, which should approach the ideal state as nearly as possible, the powder particles finally have approximately "mono particle" coatings of AEROSIL (see Figures 4 and 5, compare 2.2).

Had the method of mixing been reversed, the small amount of AEROSIL (0.5 to 1%) would have been coated or "swallowed" by the powder and the powder would have scarcely changed its flow behavior. In addition to the method of mixing, the process of mixing requires some attention: the powder component should be loosely mixed without too much pressure or shearing stress.
Figure 1. Electronmicrograph of AEROSIL 200

Figure 2. Particle size distribution of AEROSIL 200

Figure 3. Left test: NaHCO₃ untreated; right test: NaHCO₃ treated with 1% AEROSIL R972
For carrying out this type of mixing experiment, we found the "turbula" mixing for example to be quite useful. It was also sufficiently effective in producing so called "dry water" (see Figures 6 to 8).

![Figure 4. NaHCO₃ (without AEROSIL R 972)](image)

![Figure 5. NaHCO₃ + 0.5% AEROSIL R 972)](image)

5. Methods of measurement

The evaluation of a powder succeeds according to several points of view of which the following tests are the most important for practical concerns:

5.1 Ability to flow
Figure 6. "Turbula" mixer  Manufacturer: W. A. Bachofen, Basel (Switzerland)

Figure 7. Left: before mixing; right: after mixing

Figure 8. "dry water" consisting of 90% water and 10% AEROSIL R 972
5.1 Ability to flow

Several methods have been proposed for measuring the ability of a powder to flow. These are based on running the powder through standardized funnels, measurement of the angle of inclination or elevation of the powder which runs out, or on the determination of the frictional resistance of the particles to each other.

Because the measurement of the "ability to flow" is ultimately concerned with "flowing" the simple method of viscosity measurement for fluids obviously suggested itself for the measurement of the flow of powders. This method employs vessels with different sizes of mouths to measure the viscosity.

For this purpose hour glass shaped glass vessels with different sized mouths were suggested and were chosen (see Figures 9 and 10). The vessels were made hydrophobic by rinsing
them with a silicon oil solution and leaving them in a 300°C drying cupboard for 6 hours.

5.2 Dispersion behavior

Apart from an ability to flow, a powder must have a good dispersion behavior for a number of applications. This is particularly true for all powdered materials which are sprayed or scattered (cosmetic powder, pest control agents, etc.). Because these materials are so finely divided and have such a low loose weight, their flow behavior can only be improved in unusual cases. This is also not at all necessary for this application assuming the dispersion behavior to be sufficiently good.

However, because dispersion behavior is very strongly influenced by humidity (storage, transport), the following weathering test was chosen to examine it:

The powder to be tested was weathered in an open weighing bottle under 65 or 80% relative humidity at 30°C. Its external behavior was periodically checked. (Insofar as it is of interest, the flow behavior can be tested at the same time, using the method described in 5.1) (Figures 11 to 14).

5.3 Resistance to pressure

Powdered products tend to agglomerate when stored in bags. The following test has proved useful in judging their "storage resistance"; the powder to be tested is placed in a cylinder of about 50 mm inner diameter to a height of about 20 mm. A 1.2 kg piston carrying an additional 2 kg of weight is placed on the powder. The duration of the pressure test can be chosen in accordance with the requirements of the powder with respect to packing and storage. From our own experience 24 hours under

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pressure was sufficient for a judgement to be made. The pressure amounted to 0.17 kg/cm² with this test arrangement. This corresponded to the weight of 10 to 12 50 kg bags placed on top of each other. After 24 hours, the 2 kg weights were removed and the hollow cylinder was carefully raised. The powder remaining under the piston was then determined. (Figures 15 and 16).

5.4 Static charge

The static charge on a powder also considerably hinders its ability to flow. W. A. Ritschel (12) reported about this in detail.

A test which is simple to carry out is the energetic rubbing of the powder against the wall of a beaker with a glass or hard rubber rod.
In this way, powders which become charged form a stubborn coating on the beaker wall and glass rod. Uncharged powders, on the other hand, show no coating (Figures 17 and 18).

The disturbance of the flow behavior of spin sintered powders by accumulation of static charge is shown in a demonstration experiment (Figure 19).

W. A. Ritschel (12) shows the influence of static charge on the grinding of a powder for the example of proxyphyllin (obstruction of milling)* (Figures 20 and 21).

*With the friendly permission of Prof. Dr. W. A. Ritschel, Saeckingen, for which we thank him here.
### TABLE 1.

<table>
<thead>
<tr>
<th>Vessel No.</th>
<th>Evaluation No.</th>
<th>north diameter (mm)</th>
<th>evaluation for smooth running through of powder</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1</td>
<td>2.5</td>
<td>very good (seldom used with powders)</td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>5</td>
<td>good</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>7</td>
<td>good on the whole</td>
</tr>
<tr>
<td></td>
<td>4</td>
<td>12</td>
<td>adequate</td>
</tr>
<tr>
<td></td>
<td>5</td>
<td>18</td>
<td>deficient</td>
</tr>
<tr>
<td></td>
<td>6</td>
<td>substandard</td>
<td>substandard (did not pass through number 5)</td>
</tr>
</tbody>
</table>

All "hour glasses" had a height of 42 mm and an inner diameter of 90 mm

### TABLE 2.

<table>
<thead>
<tr>
<th>evaluation note</th>
<th>characteristic behavior</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 = very good</td>
<td>completely unchanged and flowing excellently (smoothly flowing through hour glass #2)</td>
</tr>
<tr>
<td>2 = good</td>
<td>not adhering to the surface; practically unchanged from its pre-test state</td>
</tr>
<tr>
<td>3 = good on the whole</td>
<td>slight surface adherence; easily returned to the original dispersion</td>
</tr>
<tr>
<td>4 = adequate</td>
<td>adhering to the surface; returns to the original dispersion by rolling and shaking</td>
</tr>
<tr>
<td>5 = deficient</td>
<td>adheres; the original dispersion cannot be reached by shaking</td>
</tr>
<tr>
<td>6 = substandard</td>
<td>agglomerated into clumps and moist to the touch</td>
</tr>
</tbody>
</table>
Figure 13. After 2 days weathering
Left: \( \text{NAHCO}_3 \) (without addition) note 5;
middle: \( \text{NaHCO}_3 \) (+0.5% AEROSIL R 972) note 1;
right: \( \text{NaHCO}_3 \) (+1% Ca-stearate) note 4

Figure 14. After 4 days weathering
left: \( \text{NaHCO}_3 \) (without addition) note 6;
middle: \( \text{NaHCO}_3 \) (+0.5% AEROSIL R 972) note: 1-2;
right: \( \text{NaHCO}_3 \) (+1% Ca-stearate) note 5

Figure 15. Test arrangement for resistance to pressure

Figure 16. Pressure resistance of ammonium chloride
left: ammonium chloride without addition
right: ammonium chloride with 1% AEROSIL R 972

Figure 17. Left: PVC without addition,
right: PVC + 1% aluminum oxide C
### TABLE 3.

<table>
<thead>
<tr>
<th>Evaluation Note</th>
<th>Characteristic Behavior</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 = very good</td>
<td>completely unchanged and flowing smoothly through hourglass #2</td>
</tr>
<tr>
<td>2 = good</td>
<td>partially loosely adhering; easily returned to the original state</td>
</tr>
<tr>
<td>3 = good on the whole</td>
<td>loosely formed; can be returned to powder with slight finger pressure</td>
</tr>
<tr>
<td>4 = adequate</td>
<td>loosely agglomerated; still crumbles under finger pressure</td>
</tr>
<tr>
<td>5 = deficient</td>
<td>semi-firmly agglomerated; no longer crumbles under finger pressure</td>
</tr>
<tr>
<td>6 = substandard</td>
<td>solidly formed</td>
</tr>
</tbody>
</table>

6. **Examples of the use of AEROSIL in improving the properties of powdered materials**

6.1 Improvement of flow behavior (Figures 22 and 26)

6.2 Improvement of weathering stability of moisture sensitive material (Figures 23 and 24)

6.3 Improvement of pressure stability (stacking resistance (Figure 25))

6.4 Improvement of pressure sensitive mixtures of materials with regard to agglomeration and clumping (Figures 27 and 28)

6.5 Reduced sintering of powdered or granulated substances (Figures 29 and 30)

6.6 Loosening and disagglomeration of spray powders (Figures 32 and 33)

6.7 Conversion of fluid materials to powdered form (Fig 31)

6.8 Increase in volume of powdered materials (Figs 34 and 35)
Figure 18. Flow behavior of statically charged and uncharged PVC powder. Left: PVC powder without addition; right: PVC powder + 1% aluminum oxide C (both samples charged in the "Turbula mixer")

Figure 19. Left: PVC spin sintered powder without addition (powder shows hesitant flow and blowing out of the pressurized air through crater forming channels) right: PVC spin sintered powder with 0.2% AEROSIL R 972 and aluminum oxide C (powder flows homogeneously without blowing out of the powder. No air pressure channels!)

Figure 20. Grinding of proxphyllin without addition

Figure 21. Grinding of proxphyllin, to which 0.45% AEROSIL 200 was added
Figure 22. Effect of different flow agents on ground sulfur.
Flow agent 1: aluminum oxide C
Flow agent 2: AEROSIL 200
Flow agent 3: tricalcium phosphate
Flow agent 4: calcium silicate

Figure 23. Weathering progress of ammonium nitrate with the addition of different flow agents (65% relative humidity, 30°C).
Flow agent 1: 1% AEROSIL R 972 (mixture not colored)
Flow agent 2: 1% AEROSIL 200 (mixture not colored)
Flow agent 3: 0.5% Berlin blue (mixture colored blue)
Flow agent 4: 1% copper-I-oxide (mixture colored red)
Figure 24. Weathering progress of sodium hydrogen carbonate with and without flow agent addition (65% relative humidity, 30°C)
Flow agent 1: 0.5% sylicic acid D 17
Flow agent: 1.0% calcium stearate
Curve 3: without flow agent addition

Figure 25. Pressure resistance with increasing addition of hydrophobic sylicic acid D 17
Curve 1: ammonium phosphate
Curve 2: ammonium sulfate
Curve 3: mixture ammonium phosphate: ammonium sulfate = 3.1
Figure 25: Remark: Example for the open observed phenomenon that the pressure resistance of material mixtures is worse than that of the individual components.

Figure 26. Left: sulfur without addition, right: sulfur + 0.2% sylicic acid D 17

Figure 27. Concentrated feed, left: without addition, right: with 10% AEROSIL 200

Figure 28. Powdered milk, left: without addition, right: with 0.5% sylicic acid D 17

Figure 29. Artificial resin formed from phenol-formaldehyde, left: without addition (sample reduced for picture) right: with 1.5% AEROSIL 200 (both samples stored 7 years)
Figure 30. Paraffin geperite
Left: without addition right: treated with 2% AEROSIL (both samples store 5 years)

Figure 32. Finely ground magnesium oxide (in the normal supply state)

Figure 34. Increase in loose volume of a body powder with the addition of increasing amounts of AEROSIL 200

Figure 31. Conversion of 70% choline chloride solution to powdered form left: 70% choline chloride right: with 40% AEROSIL

Figure 33. Similar magnesium oxide mixed with 5% hydrophobic sylicic acid D 17.

Figure 35. Equal weights of powder left: without AEROSIL right: with 10% AEROSIL 200
7. Summary and prospects

The use of AEROSIL 200, AEROSIL R 972, aluminum oxide C and sylicic acid D 17 as flow agents has made broad improvements possible in the properties of powdered products.

In addition to flow and dispersion behavior, the sifting and spray behavior and the grinding and air separation (also sifting) have been substantially improved. This is particularly true for the hydrophobic type AEROSIL R 972 which, in addition to providing outstanding weathering stability, can be used in certain cases to regulate the moisture absorbancy as desired.

Powdered materials containing AEROSIL as a flow agent can more easily and quickly be mixed to homogeneity and are far less subject to the danger of a subsequent separation (paints, artificial resins, etc.).

Sprayed materials and powders function more reliably with AEROSIL. Their improved dispersion behavior allows a substantial reduction in the cost of their application because the same amount of surface can be covered with smaller amounts of treated powder than is the case with untreated powder.

The conversion of fluid or plastic materials into powdered form by means of AEROSIL makes new forms of packing, handling and application possible. The high absorbability of AEROSIL by fluids and pastes is of particular significance in maximizing the concentration of the material in the product.

The often desensitizing effect of AEROSIL or spontaneously reactive materials and material mixtures (e.g., organic peroxides) makes a reduction in the dangers of handling and transport of these materials possible.
8. LITERATURE

(1) Prevention of caking of fertilizers, Patent 950 559 11.10 1956 (BASF)


(4) Food additives, Silicon dioxide. Federal Register 27, 4914 (25.5.1962)


(6) Food additive, silicon dioxide. Federal Register 29, 1385 (28.1.1964)


(12) W. A. Ritschel. On the use of antistatics with aggregates in the pharmaceutical processing technique. Pharm. Ind. 28, 689-694 (1966)


(14) E. Wagner and H. Bruenner. AEROSIL, manufacture, properties and behavior in organic fluids. Angew. Chem. 72, 744-750 (1960)


<table>
<thead>
<tr>
<th>PHYSICO-CHEMICAL DATA</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>AEROSIL</strong></td>
</tr>
<tr>
<td>200</td>
</tr>
<tr>
<td>appearance</td>
</tr>
<tr>
<td>moisture absorbancy surface according to BET</td>
</tr>
<tr>
<td>average size of the primary particles</td>
</tr>
<tr>
<td>compressed volumes (DIN 53 194)</td>
</tr>
<tr>
<td>drying losses (DIN 53 198 method A) (2 hrs at 105°C)</td>
</tr>
<tr>
<td>heat loss *) (DIN 52911) (2 hrs at 1000°C)</td>
</tr>
<tr>
<td>pH-value (DIN 53200) (in 4% aqueous emulsion)</td>
</tr>
<tr>
<td>SiO₂*)</td>
</tr>
<tr>
<td>Al₂O₃*)</td>
</tr>
<tr>
<td>Fe₂O₃</td>
</tr>
<tr>
<td>TiO₂</td>
</tr>
<tr>
<td>HCl</td>
</tr>
<tr>
<td>grit according to Mocker (DIN 53 580)</td>
</tr>
</tbody>
</table>

*) relative to the substance which was dried for 2 hrs at 105°C.
**) in water: methanol 1:1
(table continued)....

Packing:  
AEROSIL 200: paper bags to 10 kg net weight  
cyclic acid D 17: paper bags to 25 kg net weight  
AEROSIL R 972 and  
aluminum oxide C paper bags to 5 kg net weight

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