Silicone Adhesives for Transdermal Drug Delivery Systems

Product Applications and Benefits

Efficacy. Compliance.

Advancing healthcare through material innovations
Transdermal drug delivery patches are designed to deliver a therapeutically effective amount of drug across a patient’s skin to efficiently treat targeted diseases. Transdermal patches typically involve a pressure sensitive adhesive used as a patch fixation system and/or drug matrix.

Every specific transdermal patch typically requires a number of properties that can be drug dependent like:

- Good adhesion profile for the application period
- Adequate drug loading
- Manageable patch size
- Sufficient skin flux of the drug
- Sufficient control of the delivery rate to avoid overdosing
- Depletion rate over the designed dosage period

This brochure will help you choose the right adhesive to meet your specific needs.

Silicone adhesives are currently used in a wide range of pharmaceutical applications from hormone therapy to central nervous system related pathologies.

*Dow Corning®* BIO-PSA Silicone Adhesives are chemically very stable, non-sensitizing, non-irritating and non-cytotoxic.

U.S. Food and Drug Administration Drug Master Files and Technical Files are available for products used in pharmaceutical applications.

We work with our customers to ensure that our silicone adhesives meet their regulatory and toxicology requirements. Dow Corning is ISO 9001 Certified and compliant with regulatory guidelines appropriate to your needs.

We also work with our customers to customize our solutions to their specific needs.
Dow Corning BIO-PSA Silicone Adhesives

Silicone adhesive technology for transdermal and topical drug delivery

Transdermal and topical drug delivery systems need suitable adhesives to secure the patch to the skin and insure proper drug loading, stability and release. Dow Corning has specifically designed a line of pressure sensitive adhesives, the Dow Corning® BIO-PSAs, to prepare transdermal and topical drug delivery systems.

Dow Corning BIO-PSAs are compatible with a wide range of drugs, skin permeation enhancers and other suitable excipients, in liquid or powder forms.

Dow Corning BIO-PSAs can be formulated to provide various rates of drug permeability through the skin allowing a controlled drug release to a patient over time.

Dow Corning BIO-PSAs offer excellent skin adhesion for extended periods of time, moisture resistance, and are non-skin-irritating and non-skin-sensitizing.

Dow Corning offers a variety of BIO-PSA silicone adhesives according to their:
- Compatibility with amine functional molecules
- Polarity level (silanol content)
- Tack level
- Processing technology, solvent-based or hot melt system

Chemistry overview

Dow Corning® Standard BIO-PSAs are obtained following a condensation reaction between a silanol endblocked polydimethylsiloxane (PDMS) and a silicate resin. To produce Dow Corning® Amine-Compatible BIO-PSAs, the adhesive is further reacted with trimethylsilyl in order to reduce the silanol content of the adhesive polymer. Both standard and amine-compatible Dow Corning BIO-PSAs are then diluted in the appropriate solvent, mainly ethyl acetate or n-heptane, to obtain solvent-based materials. To produce a solvent-free version of the standard BIO-PSA, having melt characteristics applicable for hot melt coating, a PDMS plasticizer is added to the adhesive obtained from the reaction between silanol endblocked PDMS and silicate resin.
Customized Pressure Sensitive Adhesives (PSAs)

*Dow Corning* BIO-PSA Silicone Adhesives are designed for flexibility through custom formulation to meet your needs. *Dow Corning*’s current product line of BIO-PSAs allows you to adjust various critical parameters which are considered during the development of a patch drug delivery system including: adhesive performance, drug compatibility and diffusion, and coating condition (solvent based or hot melt). The safety, efficacy and stability of *Dow Corning* BIO-PSAs have been demonstrated by more than 30 years of use in the healthcare industry.

Benefits of *Dow Corning* BIO-PSA Silicone Adhesives in drug delivery

- Strong adhesion with long wear characteristics
- Customized adhesion for adapting the application, skin type, level of activity and environment
- Compatibility with amine functional drug — various silicone options are available for ease of formulation with drugs
- Controlled drug diffusion rate
- Easy removal from skin
- Non-irritating and non-sensitizing over extended wear

Processing options

*Dow Corning* BIO-PSAs can be coated as a diluted form or as a hot melt material. The solvent-based adhesives are mainly available in heptane or ethyl acetate; the solvent selection is chosen based on the drug solubility and coating criteria. The hot melt adhesive is solvent-free and can be softened with heat, then cooled to a nearly flow-less state.

Both adhesive forms are generally coated onto a suitable release liner (e.g., fluoro coated liner) and then transferred to the final substrate (e.g., backing) by laminating.

Why controlled drug delivery?

![Drug administration chart](image-url)
### Summary table of Dow Corning BIO-PSAs

<table>
<thead>
<tr>
<th>Resin/Polymer Ratio</th>
<th>Silanol Content</th>
<th>Typical Solids Content</th>
<th>Solvent</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Standard BIO-PSA</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>7-4401*</td>
<td>65/35</td>
<td>High</td>
<td>60</td>
</tr>
<tr>
<td>7-4402*</td>
<td>65/35</td>
<td>High</td>
<td>60</td>
</tr>
<tr>
<td>7-4501</td>
<td>60/40</td>
<td>High</td>
<td>60</td>
</tr>
<tr>
<td>7-4502</td>
<td>60/40</td>
<td>High</td>
<td>60</td>
</tr>
<tr>
<td>7-4601</td>
<td>55/45</td>
<td>High</td>
<td>60</td>
</tr>
<tr>
<td>7-4602</td>
<td>55/45</td>
<td>High</td>
<td>60</td>
</tr>
<tr>
<td>SRS7-4501</td>
<td>60/40</td>
<td>Medium</td>
<td>70</td>
</tr>
<tr>
<td>SRS7-4502</td>
<td>60/40</td>
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<td>60</td>
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<tr>
<td>SRS7-4602</td>
<td>55/45</td>
<td>Medium</td>
<td>60</td>
</tr>
<tr>
<td>Hot Melt 7-4560</td>
<td>60/40</td>
<td>High</td>
<td>100</td>
</tr>
</tbody>
</table>

| **Amine-Compatible BIO-PSA** |                |                        |         |
| 7-4101*                 | 65/35          | Low                    | 60      | Heptane |
| 7-4102*                 | 65/35          | Low                    | 60      | Ethyl Acetate |
| 7-4201                  | 60/40          | Low                    | 70      | Heptane |
| 7-4202                  | 60/40          | Low                    | 60      | Ethyl Acetate |
| 7-4301                  | 55/45          | Low                    | 70      | Heptane |
| 7-4302                  | 55/45          | Low                    | 60      | Ethyl Acetate |

### Typical properties of Dow Corning BIO-PSAs

#### Solvent-based BIO-PSAs

<table>
<thead>
<tr>
<th>Parts resin</th>
<th>Tack level</th>
<th>Peel adhesion (g/cm)</th>
<th>Shear (kg/6.3cm²)</th>
<th>Complex viscosity at 0.01 rad/s and 30 °C (P)</th>
</tr>
</thead>
<tbody>
<tr>
<td>55</td>
<td>High</td>
<td>500</td>
<td>15</td>
<td>5.10⁶</td>
</tr>
<tr>
<td>60</td>
<td>Medium</td>
<td>700</td>
<td>16</td>
<td>5.10⁷</td>
</tr>
<tr>
<td>65*</td>
<td>Low</td>
<td>-</td>
<td>-</td>
<td>5.10⁷</td>
</tr>
</tbody>
</table>

#### Hot melt BIO-PSAs

<table>
<thead>
<tr>
<th>Parts resin</th>
<th>Tack level</th>
<th>Peel adhesion (g/cm)</th>
<th>Shear (kg/6.3cm²)</th>
<th>Complex viscosity at 0.01 rad/s and 30 °C (P)</th>
</tr>
</thead>
<tbody>
<tr>
<td>60</td>
<td>Very high</td>
<td>300</td>
<td>11</td>
<td>5.10⁵</td>
</tr>
</tbody>
</table>

*These are used in combination to optimize the adhesive performance.*
Visco-elastic properties of Dow Corning BIO-PSAs

_Dow Corning_ BIO-PSA Silicone Adhesives are visco-elastic compounds based on the “resin-in-polymer” concept. They are typically evaluated by the dynamic rheology “oscillation test” method. The rheological parameters, viscous modulus (G’’), elastic modulus (G’) and complex viscosity (Eta*) are scrutinized for performance comparison.

**Principle of adhesion/release:** When the viscous modulus is low, optimum wetting/optimum bonding occurs. Sufficient elasticity is required to break the adhesive interface and remove the device.

The increase in resin content within the adhesive polymer — from 54% to 63% — results in increased elastic modulus (G’) and viscous modulus (G’’) values over the entire frequency range. These physical characteristics are foremost to achieve lower tack and higher cohesiveness (cold flow resistance). Good adhesive performance is obtained when G’ values are low (possibly lower than G’’) at low frequency rates (e.g., 0.01 rad/s). Nevertheless, the swift increase in G’ values (preferably higher than G’’) as the frequency is increased exhibits the optimum creep resistance behavior.

Historical data show that the higher the complex viscosity (Eta*) at low frequencies (e.g., 0.01 rad/s), the more difficult it is for an adhesive to creep. Therefore, when the resin content of a _Dow Corning_ BIO-PSA is increased, the adhesive becomes less likely to creep (cold flow).


**Impact of the resin content on the adhesive rheology**

*Comparison of the loss and storage moduli*

<table>
<thead>
<tr>
<th>Resin Content</th>
<th>G’ at 54% resin</th>
<th>G’ at 57% resin</th>
<th>G’ at 60% resin</th>
<th>G’ at 63% resin</th>
</tr>
</thead>
<tbody>
<tr>
<td>54% resin</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>57% resin</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>60% resin</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>63% resin</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Impact of the resin loading on the complex viscosity**

<table>
<thead>
<tr>
<th>Resin Content</th>
<th>Eta* (P)</th>
</tr>
</thead>
<tbody>
<tr>
<td>54 parts resin</td>
<td></td>
</tr>
<tr>
<td>57 parts resin</td>
<td></td>
</tr>
<tr>
<td>60 parts resin</td>
<td></td>
</tr>
<tr>
<td>63 parts resin</td>
<td></td>
</tr>
</tbody>
</table>

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Impact of the resin content on the adhesive rheology

Comparison of the loss and storage moduli

Impact of the resin loading on the complex viscosity
Toxicology Testing and Summaries

Our toxicology group has performed descriptive toxicology studies (acute, subchronic, reproductive testing and neurotoxicity screening) as well as more specialized investigations, including medical material testing (Class I–VI), whole-body autoradiography, pharmacokinetics, metabolism, image analysis, biochemical and in vitro/in vivo dermal absorption evaluations. We also offer customized resources for your research and consulting needs.

Our toxicology group can also provide Health Summaries, study reports and/or Health Opinions based on safety and toxicology information that has already been generated.

U.S. FDA Drug Master File and Technical File Content

- GENERAL PRODUCT INFORMATION
  - Product nomenclature, description, prior registered uses
- PRODUCT MANUFACTURE
  - Manufacturer information
  - Process description and in-process control detailed
  - Starting materials specifications
- PRODUCT CHARACTERIZATION
  - Impurity profile
- PRODUCT CONTROL
  - Specifications and quality control testing in routine formulation and composition
  - Batch analysis
- PRODUCT REFERENCE STANDARDS
- PRODUCT CONTAINER CLOSURE SYSTEMS
- PRODUCT STABILITY
  - Stability protocols
  - Stability data

Regulatory Information Sheets

To help support the registration of drug products containing these adhesives, summaries of “Product Regulatory Information” can be prepared and provided. These summaries follow the International Pharmaceutical Excipient Council (IPEC) Excipient Information Package (EIP) template and include such sections as:

- General Product Information
- Manufacturing, Packaging and Release Site
- Physicochemical Information
- Regulatory Information (based on composition and manufacturing process)
- Biocompatibility Information
- Miscellaneous Product Information
Good Manufacturing Practices†

Dow Corning® BIO-PSAs are produced, tested, and packaged under strict quality control guidelines at the Healthcare Industries Materials Site (Hemlock, MI). The site is dedicated to the production of silicone materials for healthcare applications. It is registered with the United States Food and Drug Administration as a drug establishment (CFN 1816403). The site quality system for pharmaceutical excipients utilizes principles from The Joint IPEC-PQG Good Manufacturing Practices Guide for Pharmaceutical Excipients.

Examples of GMP principles included are shown below:

- Manufacturing process controls
- Contamination control (clean plant policies)
- Documentation control system/records (including batch records and labeling systems)
- Complete product/process traceability
- Complaint management and corrective action processes
- Central review, customer notification of changes
- Appropriate building, facilities, equipment practices
- Quality system audits and third-party inspections

† Dow Corning applies the appropriate level of pharmaceutical Good Manufacturing Practices for the intended healthcare application.

REFERENCES:

Contact Dow Corning
When you need innovation, Dow Corning can help. We are dedicated to meeting your needs for specialty materials, collaborative problem solving and innovation support. We have state-of-the-art application center laboratories in our facilities. Learn how we can help you at dowcorning.com/healthcare.

Your Global Connection
Dow Corning has sales offices and manufacturing facilities worldwide, as well as full-service, global technical support. Contact us today by visiting dowcorning.com/ContactUs.