



I. Intended use

MagSi-Tools are surface activated magnetic particles, intended for covalent immobilization of proteins (e.g. antibodies, enzymes), peptides, nucleic acids or other molecules of interest. Different surface modifications and bead sizes allow for choosing the optimal product for the right molecule to be coupled, and for the intended application. Please take into consideration which groups are available on the ligand for coupling, and try to prevent inactivation or hiding the active or exposed site of the ligand.

After coupling the molecule of interest (ligand) is coupled to the magnetic particles, the resulting beads can be used in downstream applications such as:

- Isolating specific target proteins, antibodies, nucleic acids, cells, viruses, etc. (preparative applications)

- Detecting specific target proteins, nucleic acids, cells, viruses, etc. (diagnostic applications)

- Immobilizing enzymes, thereby enhancing stability and minimizing auto-catalysis. Magnetic collection of the particle/enzyme complex allows to remove the enzyme from the reaction, and to reuse it in a new reaction.

II. Principle

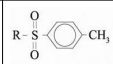
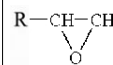
Magnetic beads are an ideal tool for immobilizing molecules (proteins, enzymes, antibodies, peptides, nucleic acids, etc.) on a solid phase, to be used for e.g. detecting, enriching, or cleaving specific target molecules. The easy and efficient collection of beads in magnetic fields allows for easy rinsing and removal of excess reagents and ligand after coupling the ligand molecule, as well as easy use in downstream applications. The use of magnetic beads does not require columns or centrifugation steps, and are therefore ideal in high-throughput and automated applications.

Selection of your MagSi-Tool particle:

Bead surfaces

MagSi-Tools are magnetic silica beads with different surface activations to best suit your needs. Surfaces available are:

Table 1: Active surfaces and example applications of MagSi-tools

| Surface activated | Formula | Example Applications |
|---|---|--|
| Silica (stored in 0.05% sodium azide) | Si-OH | - End-users' own application (e.g. functionalization of the MagSi beads) |
| Carboxyl (stored in PBS, 0.05% sodium azide) | R-COOH | - Protein and peptide immobilization - Antibody immobilization |
| Aldehyde (stored in PBS, 0.05% sodium azide) | R-CHO* | - Protein immobilization |
| Amine (stored in 0.05% sodium azide) | R-NH ₂ | - Protein immobilization |
| Sulfhydryl (stored in PBS, 0.05% sodium azide) | R-SH* | - Immobilization via target cysteine groups, coupling to gold surfaces |
| Tosyl (stored in DMSO:THF 1:1) | R-CO-N ₂ H ₂ | - Antibody immobilization - Protein and peptide immobilization |
| Hydrazide (stored in PBS, 0.05% sodium azide) |  | - Glycoprotein immobilization - Protein and peptide immobilization |
| Epoxy (stored in DMSO:THF 1:1) |  | - Enzyme immobilization - Protein and peptide immobilization |

* coupling of other organic molecules, such as nucleic acids or carbohydrates, is also possible. CHO- and SH-beads have a limited stability, and must be used for coupling ligand within 2-3 weeks after production.

Bead size

MagSi-Tools magnetic beads come in three sizes, 600 nm, 1 µm and 3 µm. 600 nm beads have the advantage of having a larger

surface area and the sedimentation time of 600nm MagSi beads is approximately 4 times slower than that of 1.0µm beads. This allows longer incubation times without shaking/mixing, and may be important in automated and other high-throughput applications in which shaking/mixing options are often lacking. MagSi beads with a diameter of 3µm have stronger magnetic properties and will separate approximately 4x faster than 600nm beads under same conditions; approximate separation time is ≤1 minute using a suitable magnet.

III. Material Supplied

- 2, 10, or 100 ml MagSi-Tools 600, 1.0 or 3.0 (supplied at 10 mg/ml).

Additional materials needed

- Buffers and Materials (depending on the application, contact for support)
- Magnetic separator for bead separation/collecting (see order information)
- Mixer/vortex to homogenize samples and resuspend beads (depending on the application, contact for support)

IV. Product usage

The products are stable at least 1 year after purchasing date when stored at 2-8°C (except CHO- and SH-beads: limited stability, must be used for coupling ligand within 2-3 weeks after production), unless mentioned otherwise on the label. Store beads in well closed vial and in upright position to prevent drying of the beads since this makes them more difficult to re-suspend. Do not freeze the product! Vortex bead suspension well before use. If you expect iron interference in downstream applications, we strongly advise you to rinse the beads before usage.

MagSi-Tools are suspended in PBS buffer or water with 0.05% sodium azide (toxic) added as a preservative, or in a 1:1 mixture of DMSO and THF. MSDS of our products can be found at our site (www.amsbio.com). Before using the beads it is important to rinse with

water or PBS to remove any components that could interfere with results.

IV. Protocols for ligand immobilization

Table 2: Coupling chemistries and conditions for different MagSi-Tools

| Bead Surface | Chemicals needed | Protein binding | Treatment | Comments |
|---------------------------------------|-----------------------------|---|---|---|
| Carboxyl ¹ (COOH) | EDC/NHS | Amine groups (from lysine and/or as unblocked N-termini) Lysine, histidine, cysteine, tyrosine etc. | No treatment needed | Can be used to couple most proteins |
| Aldehyde (CHO) | Aldehyde/Amine reaction | Amine groups | No treatment needed | Add reducing agent to stabilize amide bond |
| Thiol (SH) | Redox reaction ³ | Free cysteine | Reduce disulphides under non-denaturing conditions to generate free cysteine. | Useful for proteins containing cysteines. Risk of multiple coupling |
| Amine ² (NH ₂) | Gluteraldehyde | Amine/aldehyde | No treatment needed | Add reducing agent to stabilize amide bond |
| Tosyl | None | Sulfhydryl, Amine groups | No treatment needed | Useful for antibodies |
| Hydrazide | Sodium periodate | Oligosaccharide moieties | Oxidize glycoprotein under non-denaturing conditions. | Useful for glycoproteins |
| Epoxy | Adsorption/reaction support | Lysine, histidine, cysteine, tyrosine etc. | No treatment needed | Useful for enzymes |

¹ The first step is to activate the functional groups with N-hydroxysuccinimide in order of creating a highly reactive succinimide ester which reacts with amine groups contained in protein.

² Gluteraldehyde gives more stable protein binding than the carbodiimide reagents used with carboxylate beads.

Abbreviations: EDC, N-ethyl-N'-(dimethylaminopropyl) carbodiimide; NHS, N-hydroxysuccinimide.

³ Reduction of disulfides with 0.1 M DTE (dithioerythrol); coupling of protein at pH below isoelectric point; deactivate excess thiol with 20 mM PDEA (2-(2-pyridinyldithio) ethane-amine)/ 1M NaCl, pH 4,3

Disclaimer

For R&D use only. Not for drug, household or other uses. Products contain 0.05% sodium azide which is toxic. Avoid contact with the suspension buffer. When disposing the suspension buffer, flush with large amounts of water. Material Data Sheet (MSDS) is available on our website at www.amsbio.com.

VI. Technical Data

Table 3: Specifications of MagSi-Tools

| Product Name | MagSi-Tools | | |
|-------------------------|--|--------------------------|-----------------------|
| | 600 | 1.0 | 3.0 |
| Size | 600 nm | 1.0 µm | 3.0 µm |
| Concentration | 10 mg/ml | | |
| | beads/ml | | |
| | 8 - 20 · 10 ⁹ | 6 - 12 · 10 ⁹ | 1-3 · 10 ⁹ |
| Supplied product volume | 2 ml, 10 ml, 100 ml | | |
| Material | Magnetic silica beads with activated surface | | |
| Size Distribution | D5-D95 | | |
| | 500 - 900 nm | 0.7 - 1.4 µm | 0.6-7.0µm |
| Sedimentation | | | |
| | MagSi-Tools, surface activated: PBS (pH 7.4), 0.05% sodium azide (NaN ₃ , Toxic!), except: 1) MagSi-S, unmodified silica beads and MagSi-NH ₂ , amine-modified silica beads: water, 0.05% sodium azide 2) epoxy- and tosyl-activated beads are supplied in DSMO:THF 1:1. | | |
| Solution additives | | | |
| Storage | Store at 2-8°C | | |

VII. Additional Information

Order Information

| Product name | Volume | Art. No. | Product name | Volume | Art. No. |
|-----------------------------|----------------------|-------------------------------|-----------------------|----------------------|-------------------------------|
| MagSi-S 600 | 2ml 10ml 100ml | MD16003 MD18003 MD19003 | MagSi-S CHO 600 | 10ml 100ml | MD18007 MD19007 |
| MagSi-S 1.0 | 2ml 10ml 100ml | MD01003 MD03003 MD04003 | MagSi-S CHO 1.0 | 10ml 100ml | MD03007 MD04007 |
| MagSi-S 3.0 | 2ml 10ml 100ml | MD41003 MD43003 MD44003 | MagSi-S CHO 3.0 | 10ml 100ml | MD43007 MD44007 |
| MagSi-S COOH 600 | 2ml 10ml 100ml | MD16004 MD18004 MD19004 | MagSi-S Tosyl 600 | 2ml 10ml 100ml | MD16008 MD18008 MD19008 |
| MagSi-S COOH 1.0 | 2ml 10ml 100ml | MD01004 MD03004 MD04004 | MagSi-S Tosyl 1.0 | 2ml 10ml 100ml | MD01008 MD03008 MD04008 |
| MagSi-S COOH 3.0 | 2ml 10ml 100ml | MD41004 MD43004 MD44004 | MagSi-S Tosyl 3.0 | 2ml 10ml 100ml | MD41008 MD43008 MD44008 |
| MagSi-S NH ₂ 600 | 2ml 10ml 100ml | MD16005 MD18005 MD19005 | MagSi-S Hydrazide 600 | 2ml 10ml 100ml | MD16013 MD18013 MD19013 |
| MagSi-S NH ₂ 1.0 | 2ml 10ml 100ml | MD01005 MD03005 MD04005 | MagSi-S Hydrazide 1.0 | 2ml 10ml 100ml | MD01013 MD03013 MD04013 |
| MagSi-S NH ₂ 3.0 | 2ml 10ml 100ml | MD41005 MD43005 MD44005 | MagSi-S Hydrazide 3.0 | 2ml 10ml 100ml | MD41013 MD43013 MD44013 |
| MagSi-S SH 600 | 10ml 100ml | MD18006 MD19006 | MagSi-S Epoxy 600 | 2ml 10ml 100ml | MD16010 MD18010 MD19010 |
| MagSi-S SH 1.0 | 10ml 100ml | MD03006 MD04006 | MagSi-S Epoxy 1.0 | 2ml 10ml 100ml | MD01010 MD03010 MD04010 |
| MagSi-S SH 3.0 | 10ml 100ml | MD43006 MD44006 | MagSi-S Epoxy 3.0 | 2ml 10ml 100ml | MD44010 MD44010 MD44010 |