## Development of third-generation antibody-drug conjugate linker technology capable of site-selectively conjugating payloads without antibody modification

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(A) AbTis

| ONCOLOGY                 | ead  |
|--------------------------|--|
| Product Type             | Small Molecules, Antibody Drug Conjugate(ADC)  |
| Indication               | Target antigen-positive gastric cancer, breast cancer, lung cancer and pancreatic cancer etc.  |
| Target                   | Antigen present on the surface of cancer cells<br>(Ex. HER2, HER3, Nectin-4, Claudin 18.2, Trop-2, EGFR etc.)  |
| MoA(Mechanism of Action) | Binding to target antigen $\rightarrow$ Internalization $\rightarrow$ Cleaved and released cytotoxic payload in lysosome $\rightarrow$ Cancer cell death by released payload   |
| Competitiveness          | <ul> <li>Site-selective ADC obtained with controllable DAR and improved homogeneity</li> <li>Native antibody can be used without need for engineering or glycan remodeling</li> <li>Stable isopeptide bond formed between protein and linker</li> <li>Simple process 2-step process with high yield</li> <li>Applicable to different IgG subtypes</li> </ul> |
| Development Stage        | Lead   |
| Route of Administration  | IV infusion  |

