

TRODELVY®

sacituzumab govitecan (*Sah see TOO zoo mab GOH vih TEE kan*)

Consumer Medicine Information

This leaflet provides important information about using TRODELVY. **You should also speak to your doctor or pharmacist if you would like further information or if you have any concerns or questions about using TRODELVY.**

Where to find information in this leaflet:

1. Why am I being given TRODELVY?
2. What should I know before I receive TRODELVY?
3. What if I am taking other medicines?
4. How am I given TRODELVY?
5. What should I know while receiving TRODELVY?
6. Are there any side effects?
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1. Why am I being given TRODELVY?

TRODELVY contains the active ingredient sacituzumab govitecan.

TRODELVY belongs to a group of medicines known as anti-neoplastic (anti-cancer) agents. There are different classes of anti-cancer agents.

TRODELVY is designed to work differently to traditional anti-cancer agents (e.g., chemotherapy).

TRODELVY is made up of three components:

- a monoclonal antibody that recognises and attaches to certain cancer cells, and
- a substance intended to kill cancer cells, and
- a linker that links the antibody to the substance.

TRODELVY is designed to target and deliver the anti-cancer substance directly to certain cancer cells to stop the growth and spread of the cancer.

TRODELVY is used to treat adults with a type of breast cancer that:

- is oestrogen and progesterone hormone receptor (HR) negative, and human epidermal growth factor receptor 2 (HER2) negative (also called triple-negative breast cancer), and
- has spread to other parts of the body or cannot be removed by surgery, and
- has previously been treated with at least two therapies for breast cancer.