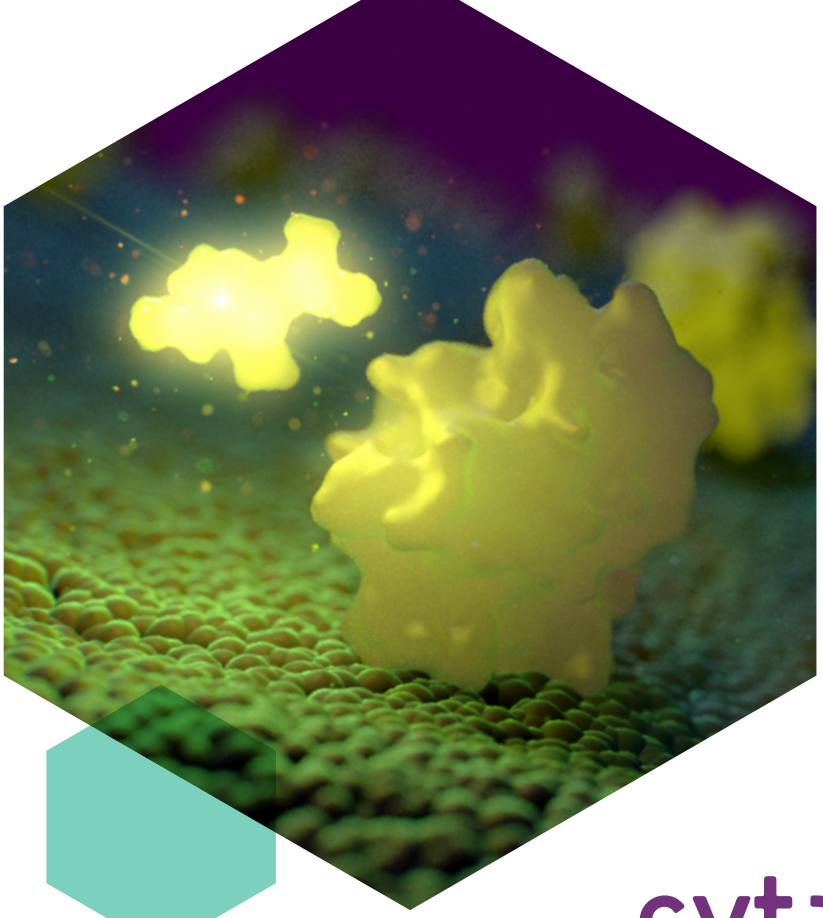

*For illustrative purposes only

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See Cancer in Real Time

CYTALUX® is the only intraoperative molecular imaging agent that makes ovarian cancer visible in real time.

INDICATION AND USAGE

CYTALUX® is an FDA approved optical imaging agent indicated as an adjunct for intraoperative identification of:

- Malignant lesions in adult patients with ovarian cancer.
- Malignant and non-malignant pulmonary lesions in adult patients with known or suspected cancer in the lung.

IMPORTANT SAFETY INFORMATION

RISK OF MISINTERPRETATION

Errors may occur with the use of CYTALUX during intraoperative fluorescence imaging to detect ovarian cancer and lesions in the lung, including false negatives and false positives. Non-fluorescing tissue in the surgical field does not rule out the presence of ovarian cancer or lesions in the lung. Fluorescence may be seen in normal tissues including bowel, kidneys, lymph nodes, lungs, and inflamed tissue.

Please see additional Important Safety Information throughout and accompanying full Prescribing Information.

ONCTARGET LABORATORIES

A Complete Resection is the Goal of Treatment for Ovarian Cancer

The amount of gross residual disease after cytoreductive surgery in ovarian cancer patients has been demonstrated to be an important independent predictor of clinical outcomes.[1] However, debulking surgery in women with advanced stage ovarian cancer often results in suboptimal outcomes.

> Studies show that among patients reported to have undergone optimal cytoreduction, 40% were found to have measurable disease on 30-day postoperative imaging.[2]

Advancements in Technology

Currently, visual inspection and palpation during surgery are the main tools utilized to estimate the extent of tumor involvement and to guide debulking. Fluorescence-Guided Surgery is an imaging technique that uses fluorescent dye to identify anatomic structures during surgical procedures using a near-infrared imaging system.

> Newer, more targeted technologies are being developed to improve intraoperative visualization of cancerous tissue.

MECHANISM OF ACTION

CYTALUX®: The First Targeted Intraoperative Molecular Imaging Agent That Illuminates Ovarian Cancer In Real Time

FOLATE

- Folate is an essential vitamin **required for cell growth** and DNA replication.[3]
- Rapidly dividing cancer cells requires an abundant supply of folate, therefore, it is **consumed in elevated quantities.**[3]
- Most ovarian cancers overexpress high-affinity folate receptors to **increase folate uptake for tumor growth**.[4]

HOW CYTALUX WORKS

CYTALUX is a small molecule that circulates quickly through the body. → **CYTALUX binds to the folate receptors on cells.** → **CYTALUX is endocytosed and concentrates in folate receptor positive (FR+) ovarian cancer tissues.** → **CYTALUX contains a bright fluorophore which is illuminated via near-infrared light.**

It can be infused intravenously in as little as an hour before surgery.

It is targeted, preferentially binding to FR+ ovarian cancer cells.

Endocytosis allows it to be concentrated in the tissues throughout the surgical procedure.

Imaging under near-infrared light enhances intraoperative visualization in challenging environments and below the tissue surface.[5]

IMPORTANT SAFETY INFORMATION

INFUSION-RELATED REACTIONS

Adverse reactions including nausea, vomiting, abdominal pain, flushing, hypersensitivity, elevation in blood pressure, dyspepsia, and chest discomfort were reported in patients receiving CYTALUX in clinical studies. A total of 17% of patients experienced reactions during administration of CYTALUX. Reactions typically occurred within 15 minutes of the start of infusion. CYTALUX infusion interruption or discontinuation due to adverse reactions occurred in 11% of all patients. Treatment with antihistamines and/or anti-nausea medication may be used. If an adverse reaction occurs during administration, the infusion can be interrupted and resumed after treatment of the reaction.

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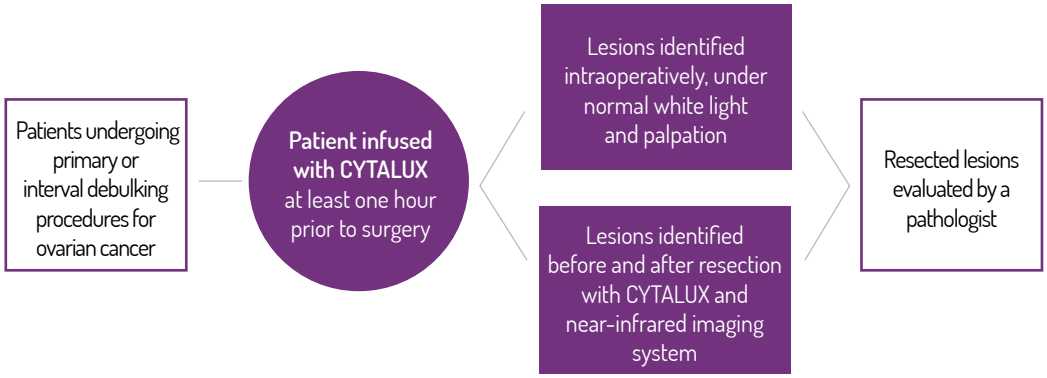
CYTALUX® Detected Additional Ovarian Cancer which would have been left behind

PHASE 3 (006 STUDY): CYTALUX FOR FR+ OVARIAN CANCER

A Phase 3, Randomized, Single Dose, Open-Label Study to Investigate the Safety and Efficacy of CYTALUX for Intraoperative Imaging of Folate Receptor Positive Ovarian Cancer



Primary Endpoint:
Proportion of patients with at least one evaluable FR+ ovarian cancer lesion confirmed by central pathology that was detected using the combination of CYTALUX and fluorescent light but not under normal light or palpation.

N=150 patients infused with CYTALUX; N=134 patients analyzed for primary and secondary endpoints

WITH CYTALUX, ADDITIONAL LESIONS WERE FOUND IN 27% OF PATIENTS*

* Women highly suspicious for or with confirmed ovarian cancer who underwent both normal and fluorescent light evaluation (Intent-to-Image set); N=134, 95% CI [19.6, 35.2]

In a subgroup analysis of patients with confirmed FR+ ovarian cancer who underwent interval debulking surgery

ADDITIONAL LESIONS WERE FOUND IN 40% OF INTERVAL DEBULKING PATIENTS**

** Phase 3 (006 Study): CYTALUX FOR FR+ OVARIAN CANCER; N=58, 95% CI [27.0, 53.4]
This subgroup analysis utilized a smaller analysis set than the primary endpoint and was not adjusted to control for error, so the results are not conclusive and should be interpreted cautiously.

Patient-level false positive rate with respect to the detection of ovarian cancer lesions confirmed by central pathology was 20%; 95% CI [0.137, 0.280]

Drug-Related Adverse Events

Phase 3 (006 Study): CYTALUX FOR FR+ OVARIAN CANCER N=150 study subjects infused with CYTALUX

DRUG-RELATED ADVERSE EVENT	MILD/MODERATE n (%)	SEVERE n (%)
Subjects with at least one drug-related TEAE	43 (28.7%)	2 (1.3%)
Total number of drug-related TEAEs	63	2

MOST COMMON DRUG-RELATED ADVERSE EVENTS	MILD/MODERATE n (%)	SEVERE n (%)
Nausea	27 (18.0%)	0
Vomiting	8 (5.3%)	0
Abdominal Pain	7 (4.7%)	0

ZERO drug-related serious adverse events were observed in any patient dosed with CYTALUX.

97% of all drug-related adverse events were mild-to-moderate.

Workflow Integration

CYTALUX® seamlessly integrates in the perioperative surgical process

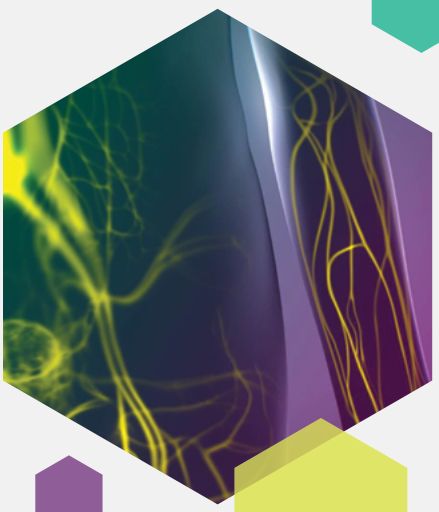
PREPARE
the recommended dose of CYTALUX as a single intravenous infusion.

ADMINISTER
CYTALUX during a 60-minute infusion, 1 to 9 hours prior to surgery for ovarian cancer.

ILLUMINATE
ovarian cancer lesions intraoperatively using a near-infrared imaging system cleared for use with CYTALUX.

A near-infrared imaging system is used to illuminate CYTALUX

- Clinical data demonstrates NIR imaging devices which excite at 760nm to 785 nm and detect emission at 790 nm to 815 nm are suitable for use with CYTALUX.
- CYTALUX is to be used with a near-infrared (NIR) imaging system cleared by FDA for specific use with pafolacianine.
- CYTALUX should only be used by surgeons who have completed a training program on the use of NIR imaging systems for fluorescence imaging during surgery. Training is provided by the device manufacturer.



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IMPORTANT SAFETY INFORMATION

EMBRYO-FETAL TOXICITY

Based on its mechanism of action, CYTALUX may cause fetal harm when administered to a pregnant woman. Advise females of reproductive potential of the potential risk to a fetus. Verify pregnancy status of females of reproductive potential prior to initiating CYTALUX treatment.

RISK OF PAFOLACIANINE AGGREGATION AND INFUSION REACTIONS

Use of the incorrect diluent to prepare the CYTALUX infusion solution can cause the aggregation of pafolacianine; aggregation may induce infusion reactions such as nausea, vomiting, abdominal pain or rash. Use only 5% Dextrose Injection to prepare the CYTALUX infusion solution. Do not use other diluents.

Please see additional Important Safety Information throughout and accompanying full Prescribing Information.

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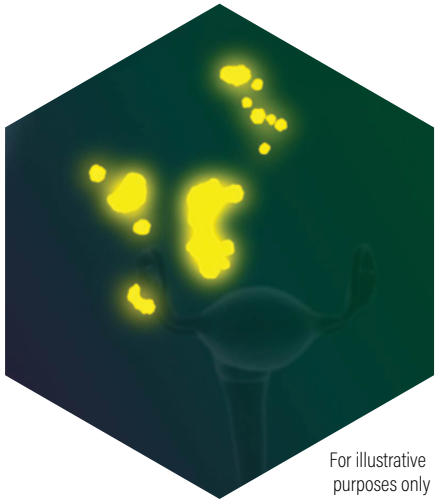

For illustrative purposes only

The use of CYTALUX helped surgeons detect additional cancer which would have otherwise been left behind in 27% of patients.*

*Women highly suspicious for or with confirmed ovarian cancer who underwent both normal and fluorescent light evaluation (Intent-to-Image set); N=134, 95% CI [0.196, 0.352]

CYTALUX® Gives You An Advantage

CYTALUX is the first targeted intraoperative molecular imaging agent that illuminates ovarian cancer in real time, enabling the detection of more cancer for resection.

Folate receptors are overexpressed in most ovarian cancers.[4]

CYTALUX binds to folate receptors and is fluoresced intraoperatively using a near-infrared imaging system.

Surgeons indicated the use of CYTALUX **altered their surgical plan in 54% of their cases.****

** Phase 3 (006 Study): Based on a prespecified exploratory endpoint for the proportion of subjects for whom the fluorescence surgical plan was changed based on fluorescence imaging both prior to initiation of the surgical procedure and upon re-imaging of the surgical field after surgical procedure prior to closing. (N=109, 95% CI [44.3,63.7])

INDICATION AND USAGE

CYTALUX® is an FDA approved optical imaging agent indicated as an adjunct for intraoperative identification of:

- Malignant lesions in adult patients with ovarian cancer.
- Malignant and non-malignant pulmonary lesions in adult patients with known or suspected cancer in the lung.

IMPORTANT SAFETY INFORMATION

ADVERSE REACTIONS

Adverse reactions that occurred in > 1% of patients were: nausea (13%), vomiting (5%), abdominal pain (2%), flushing (2%), other infusion-related reactions (2%), hypersensitivity (2%), elevation in blood pressure (1%), dyspepsia (1%), and chest discomfort (1%).

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[1] Chang SJ, Hodeib M, Chang J, Bristow RE. Survival impact of complete cytoreduction to no gross residual disease for advanced-stage ovarian cancer: a meta-analysis. Gynecol Oncol. 2013 Sep;130(3):493-8. doi: 10.1016/j.ygyno.2013.05.040. Epub 2013 Jun 6. PMID: 23747291.

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