

Boost lung tumor necrosis

EmboCept[®] S DSM 50 µm: Degradable starch microspheres for temporary transpulmonary chemoembolization (TPCE)





>60%

of patients present with late-stage III-IV lung cancer at the time of diagnosis¹

Which treatment options remain?



options for patients with late-stage

lung cancer

Who is eligible for DSM-TPCE?

- Patients in good general condition
- Without or with only minor cardiovascular comorbidities
- With sufficient lung function
- Non-thrombosed A. pulmonalis

DSM 50 µm: UNLOCKING THE BENEFITS OF TPCE

Thanks to excellent tolerability and high efficacy, transpulmonary chemoembolization (TPCE) with DSM is **easy on the patient** but **tough on the tumor**.



Figure 1: Tumor volume differences before and after treatment in a solitary metastasis rat model of CC531 adenocarcinoma (modified from Schneider et al. 2002)⁵

High flexibility

EmboCept® S DSM 50 µm can be combined with various chemotherapeutic agents⁶

Temporary embolization

Half-life of 30-40 minutes⁷

Precisely calibrated

At least 95% of microspheres are between $20-90\,\mu\text{m},$ with $50{\pm}7~\mu\text{m}$ mean size^7

HOW TO PERFORM TPCE WITH DEGRADABLE STARCH MICROSPHERES



1. Preinterventional evaluation

Control of lab parameters, clinical status, and CT/MRI scans



2. Regional anesthesia

Application of 1% mepivacain via 7F sheath into right femoral vein



3. Catheter insertion

Insertion of 5F headhunter catheter into left or right pulmonary artery via transvenous access



4. Angiography

Injection of 20 ml of contrast medium to survey arterial system



5. Balloon catheter (optional)

Insertion of catheter (diameter: 6–8 mm, length: 100–300 mm) into segmental pulmonary artery



6. Catheter advancement

Using guidewire, catheter is advanced further into subsegmental pulmonary arteries



7. Angiography

Contrast-enhanced angiographic series (with catheter blocked) for detection of arteriovenous shunts

TPCE steps based on Vogl et al. 2008⁸



8. Chemoembolization

To achieve blood flow stasis: injection of chemotherapeutic agent mixed with DSM under fluoroscopic guidance

DSM-TPCE can be combined with various chemotherapeutic agents^{7,8,9,10}:

- Mitomycin C
- Cisplatin
- Gemcitabine
- Irinotecan



9. Pressure dressing

Application following removal of catheters



10. Postinterventional evaluation

Control of lab parameters, clinical status, and CT/MRI scans



11. Repetition of treatment

At least 2–3 rounds of treatment, with intervals of four weeks

Figure 2: Illustration of the TPCE technique, depicting the insertion of the catheter via the right femoral vein into the pulmonary artery

THE CASE FOR DSM 50 µm: CLINICAL EVIDENCE

PROVEN **SAFETY**

Well tolerated

- No major complications^{2,8,9,10,11}
- No pulmonary hemorrhage, cardiac failure, or pneumothorax⁹
- No non-target embolization, e.g., to the brain⁹

Reversible DSM-TPCE

• 30–40 minutes half-life⁷

Example: effects of temporary embolization

Prior to embolization





Angiography showing the effects of temporary embolization in a patient with lung metastases. (a) Prior to embolization. (b) After embolization. Images courtesy of T. Vogl, Frankfurt/M., Germany.

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PROVEN EFFICACY

Primary lung tumors



>60%

of patients (n=17) with primary lung tumors showed a tumor volume reduction or stable disease after DSM-TPCE



Figure 3: Tumor response according to the definition of the World Health Organization following DSM-TPCE in patients with primary lung cancer (modified from Vogl et al. 2007)²

Response = volume reduction of >25% Stable disease = nonsignificant volume change Progression = volume increase of >10%

Lung metastases



Figure 4: Tumor response according to RECIST following DSM-TPCE in patients with lung metastases of different origin (modified from Vogl et al. 2008)⁸



References

- Osmani L et al. (2018): Current WHO guidelines and the critical role of immunohistochemical markers in the subclassification of non-small cell lung carcinoma (NSCLC): Moving from targeted therapy to immunotherapy. Semin Cancer Biol 52(1): 103–109
- Vogl TJ et al. (2007): Transpulmonale Chemoembolisation (TPCE) als palliatives Behandlungskonzept bei primären Lungentumoren. Fortschr Röntgenstr 179(03): 300–307; (concerns Spherex[®])
- Peters S et al. (2017): Phase II Trial of Atezolizumab As First-Line or Subsequent Therapy for Patients With Programmed Death-Ligand 1-Selected Advanced Non-Small-Cell Lung Cancer (BIRCH). J Clin Oncol 35: 2781–2789
- Genestreti G et al. (2014): Third- and further-line therapy in advanced non-small-cell lung cancer patients: an overview. Future Oncol 10(13): 2081–2096
- Schneider P et al. (2002): Chemoembolization of the lung improves tumor control in a rat model. Clin Cancer Res 8(7): 2463–2468; (concerns Spherex[®])

- 6. EmboCept® S DSM 50 μm instructions for use. Date of information: 15.05.2020
- 7. Data on file
- VogITJ et al. (2008): Transpulmonary chemoembolization (TPCE) as a treatment for unresectable lung metastases. Eur Radiol 18(11): 2449–2455; (concerns Spherex[®])
- Vogl TJ et al. (2016): Feasibility of assessing pulmonary blood volume using C-arm CT during transpulmonary chemoperfusion and chemoembolization in primary and secondary lung tumours. Br J Radiol 89(1062): 20150244; (concerns EmboCept® S)
- Vogl TJ et al. (2018): Transvenous pulmonary chemoembolization (TPCE) for palliative or neoadjuvant treatment of lung metastases. Eur Radiol 29(4): 1939–1949; (concerns EmboCept® S)
- Vogl TJ et al. (2005): Treatment of unresectable lung metastases with transpulmonary chemoembolization: preliminary experience. Radiology 234(3): 917–922; (concerns Spherex[®])







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