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through the skin — a procedure called “percutaneous ablation” — is one way to treat these metastases in the liver.

Computed tomography(CT) scanning has been used to evaluate the ablated area after treatment. But CT scanning cannot always distinguish between new cancer cells and healthy liver cells that have been affected by the ablation.

With the new approach, liver tissue removed during the ablation procedure is examined in a laboratory to see if researchers can identify any markers that predict the effectiveness of the ablation and how well patients will fare. Patients will be followed for three years after the procedure.

Who Can Join

To be eligible for this study, patients must meet several criteria, including but not limited to the following:

Patients must have liver metastases (5 cm or less in diameter).

Patients must have percutaneous liver ablation instead of surgery as a form of treatment.

Patients must be age 18 or older.

For more information and to inquire about eligibility for this study, please contact [Dr. Constantinos Sofocleous](#) at [212-639-3379](tel:212-639-3379).

Protocol

09-122

Investigator

[Constantinos T. Sofocleous](#)

Katia Manova-Todorova

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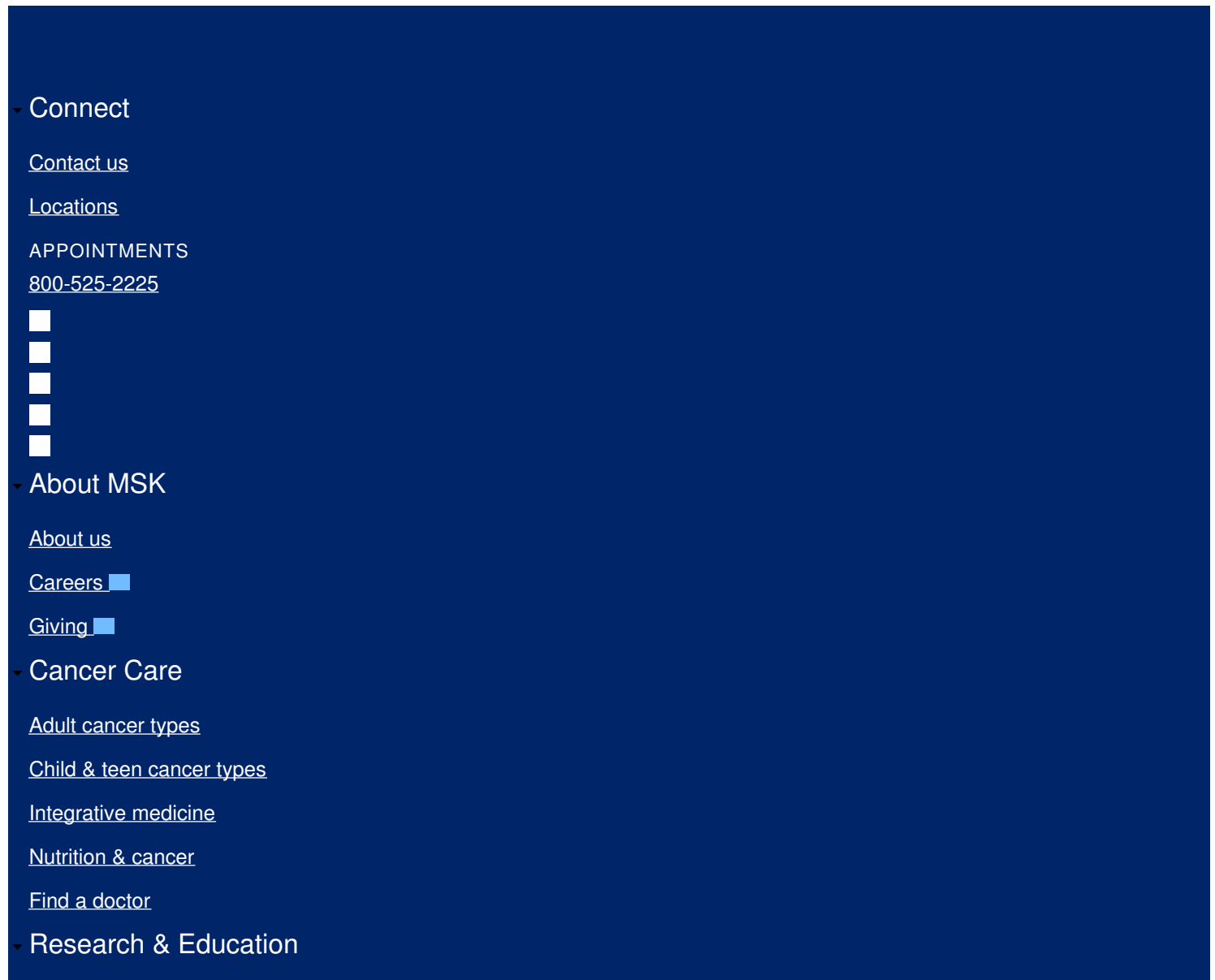
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ClinicalTrials.gov ID

NCT01494324

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