

ICMJE DISCLOSURE FORM

Date: 4/22/22

Your Name: Saba Shafi

Manuscript Title: Sporadic Mesenteric Desmoid-Type Fibromatosis with “Double-Hit” T41A and S45P Beta-Catenin Mutation Profile, A Case Report of An Extremely Rare Event: Clinically Relevant or Much Ado About Nothing?

Manuscript number (if known): PCM-22-2-CL

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Time frame: Since the initial planning of the work			
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Time frame: past 36 months			
2	Grants or contracts from any entity (if not indicated in item #1 above).	<input checked="" type="checkbox"/> None	
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11	Stock or stock options	<input checked="" type="checkbox"/> None	
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13	Other financial or non-financial interests	<input checked="" type="checkbox"/> None	

Please summarize the above conflict of interest in the following box:

None

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Date: 4/21/22

Your Name: Ashley Patton

Manuscript Title: Sporadic Mesenteric Desmoid-Type Fibromatosis with “Double-Hit” T41A and S45P Beta-Catenin Mutation Profile, A Case Report of An Extremely Rare Event: Clinically Relevant or Much Ado About Nothing?

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ICMJE DISCLOSURE FORM

Date: April 20, 2022

Your Name: Alan Rogers, MD

Manuscript Title: Sporadic Mesenteric Desmoid-Type Fibromatosis with “Double-Hit” T41A and S45P Beta-Catenin Mutation Profile, A Case Report of An Extremely Rare Event: Clinically Relevant or Much Ado About Nothing?

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ICMJE DISCLOSURE FORM

Date: 04/22/2022

Your Name: Steve Oghumu

Manuscript Title: Sporadic Mesenteric Desmoid-Type Fibromatosis with “Double-Hit” T41A and S45P Beta-Catenin Mutation Profile, A Case Report of An Extremely Rare Event: Clinically Relevant or Much Ado About Nothing?

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Date: April 22, 2022

Your Name: Valerie Grignol, MD

Manuscript Title: Sporadic Mesenteric Desmoid-Type Fibromatosis with “Double-Hit” T41A and S45P Beta-Catenin Mutation Profile, A Case Report of An Extremely Rare Event: Clinically Relevant or Much Ado About Nothing?

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Date: April 20th, 2022

Your Name: O. Hans Iwenofu, MD

Manuscript Title: Sporadic Mesenteric Desmoid-Type Fibromatosis with “Double-Hit” T41A and S45P Beta-Catenin Mutation Profile, A Case Report of An Extremely Rare Event: Clinically Relevant or Much Ado About Nothing?

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