



SDI Review Form 1.6

Journal Name:	Asian Journal of Applied Chemistry Research
Manuscript Number:	Ms_AJACR_50912
Title of the Manuscript:	Validated Stability – Indicating Methods for Determination of Sofosbuvir by UPLC and HPTLC in Pure Form and Tablet Dosage forms.
Type of the Article	Original Research Article

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This journal's peer review policy states that **NO** manuscript should be rejected only on the basis of '**lack of Novelty**', provided the manuscript is scientifically robust and technically sound. To know the complete guideline for Peer Review process, reviewers are requested to visit this link:

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PART 1: Review Comments

	Reviewer's comment	Author's comment (if agreed with reviewer, correct the manuscript and highlight that part in the manuscript. It is mandatory that authors should write his/her feedback here)																																																												
<p>Compulsory REVISION comments</p>	<p>Copy of Revision cited are attached to this document Needed</p> <p style="text-align: center;"><u>Compulsory Revision Comments</u></p> <table border="1" data-bbox="730 638 2050 1724"> <thead> <tr> <th>Page number</th> <th>Line number</th> <th>Revisions raised</th> <th>Revisions corrected</th> </tr> </thead> <tbody> <tr> <td>2</td> <td>19,20,21</td> <td>against HCV. The HCV polymerase (NS5B protein) is an RNA-dependent RNA polymerase (RdRp). It is the essential initiating and catalytic subunit of this replication complex and is critical for the viral replication cycle (2).</td> <td>for the treatment of adult patients with chronic hepatitis C virus (HCV) infection. It interferes with the HCV lifecycle, restraining viral replication.</td> </tr> <tr> <td>3</td> <td>51</td> <td>of 5N NaOH for 3.5 h</td> <td>of 5N NaOH for 3 h</td> </tr> <tr> <td>3</td> <td>59,60</td> <td>into 100- mL volumetric flask</td> <td>into 100 mL volumetric flask</td> </tr> <tr> <td>3</td> <td>61</td> <td>claimed to contain degradates derived from 0.1mg mL⁻¹ intact drug which was used for the</td> <td>claimed to contain degradates derived from 0.1mg mL⁻¹ of intact drug, which was used for the</td> </tr> <tr> <td></td> <td>72</td> <td>obtain working standard solution of 0.1mg mL⁻¹.</td> <td>obtain working standard solution of concentration 0.1mg mL⁻¹.</td> </tr> <tr> <td>4</td> <td>77,78</td> <td>Kinetex 2.6μ C₁₈ 100A (4.6-mm × 100-mm)</td> <td>Kinetex 2.6μ C₁₈ column 100A (4.6-mm × 100-mm)</td> </tr> <tr> <td>4</td> <td>80,81</td> <td>at flow rate 1 mL min⁻¹; the injection volume was 10 μL and the detection at 260 nm.</td> <td>at a flow rate 1.0 mL min⁻¹ with an injection volume of 10 μL and the detection at 260 nm.</td> </tr> <tr> <td>4</td> <td>95</td> <td>Triplicate 10 μL of each solution were injected into the UPLC system and chromatographed</td> <td>10 μL of each solution were injected into the UPLC system in triplicate and chromatographed</td> </tr> <tr> <td>4</td> <td></td> <td>Different aliquots of of intact sofosbuvir working standard solution (0.1mg mL⁻¹)</td> <td>Different aliquots of intact sofosbuvir working standard solution (0.1mg mL⁻¹)</td> </tr> <tr> <td>5</td> <td>113</td> <td>Different aliquots of stock drug solution (1 mg mL⁻¹) containing (1–11 mg)</td> <td>Different aliquots of stock drug solution (1 mg mL⁻¹) containing (1.0–11 mg)</td> </tr> <tr> <td>5</td> <td>130</td> <td>sofosbuvir in presence of its degradation product by UPLC and HPTLC methods</td> <td>sofosbuvir in presence of its degradation products by UPLC and HPTLC methods</td> </tr> <tr> <td>6</td> <td>176</td> <td>- conc. ammonia (2.5: 6: 1.5 v/v/v).</td> <td>- conc. ammonia(2.5: 6: 1.5 %v/v/v).</td> </tr> <tr> <td>13</td> <td>278</td> <td>*Avarage</td> <td>*Average</td> </tr> <tr> <td>16</td> <td>324</td> <td>1. https://www.drugbank.ca/drugs/DB08934.</td> <td>https://www.drugbank.ca/drugs/DB08934.</td> </tr> </tbody> </table>	Page number	Line number	Revisions raised	Revisions corrected	2	19,20,21	against HCV. The HCV polymerase (NS5B protein) is an RNA-dependent RNA polymerase (RdRp). It is the essential initiating and catalytic subunit of this replication complex and is critical for the viral replication cycle (2).	for the treatment of adult patients with chronic hepatitis C virus (HCV) infection. It interferes with the HCV lifecycle, restraining viral replication.	3	51	of 5N NaOH for 3.5 h	of 5N NaOH for 3 h	3	59,60	into 100- mL volumetric flask	into 100 mL volumetric flask	3	61	claimed to contain degradates derived from 0.1mg mL ⁻¹ intact drug which was used for the	claimed to contain degradates derived from 0.1mg mL ⁻¹ of intact drug, which was used for the		72	obtain working standard solution of 0.1mg mL ⁻¹ .	obtain working standard solution of concentration 0.1mg mL ⁻¹ .	4	77,78	Kinetex 2.6μ C ₁₈ 100A (4.6-mm × 100-mm)	Kinetex 2.6μ C ₁₈ column 100A (4.6-mm × 100-mm)	4	80,81	at flow rate 1 mL min ⁻¹ ; the injection volume was 10 μL and the detection at 260 nm.	at a flow rate 1.0 mL min ⁻¹ with an injection volume of 10 μL and the detection at 260 nm.	4	95	Triplicate 10 μL of each solution were injected into the UPLC system and chromatographed	10 μL of each solution were injected into the UPLC system in triplicate and chromatographed	4		Different aliquots of of intact sofosbuvir working standard solution (0.1mg mL ⁻¹)	Different aliquots of intact sofosbuvir working standard solution (0.1mg mL ⁻¹)	5	113	Different aliquots of stock drug solution (1 mg mL ⁻¹) containing (1–11 mg)	Different aliquots of stock drug solution (1 mg mL ⁻¹) containing (1.0–11 mg)	5	130	sofosbuvir in presence of its degradation product by UPLC and HPTLC methods	sofosbuvir in presence of its degradation products by UPLC and HPTLC methods	6	176	- conc. ammonia (2.5: 6: 1.5 v/v/v).	- conc. ammonia(2.5: 6: 1.5 %v/v/v).	13	278	*Avarage	*Average	16	324	1. https://www.drugbank.ca/drugs/DB08934 .	https://www.drugbank.ca/drugs/DB08934 .	
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Optional/General comments		

PART 2:

	Reviewer's comment	Author's comment <i>(if agreed with reviewer, correct the manuscript and highlight that part in the manuscript. It is mandatory that authors should write his/her feedback here)</i>
Are there ethical issues in this manuscript?	<i>(If yes, Kindly please write down the ethical issues here in details)</i>	

Reviewer Details:

Name:	D. Ramachandran
Department, University & Country	Acharya Nagarjuna University, India