



RWANDA FDA

Rwanda Food and Drugs Authority

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QMS N^o: DIS/CKL/079
 Revision No: 0
 Effective Date: 21/02/2022
 Ref. Doc: Guidelines
 DIS/GDL/005

REQUIREMENTS FOR RE – INSPECTION		
Premise name: Date :/...../.....		
Documents	YES	NO
1 Re-inspection application letter addressed to the Director General of Rwanda FDA, mentioning the proposed dates.		
2 The proof of payment of prescribed re-inspection fees		
3 A Corrective Actions and Preventive Actions (CAPA) report, detailing what has been implemented with respective visual proof and timelines for non-implemented recommendations.		

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