## Please complete as many details as possible and forward within one business day to:

Program Details													
Form Completed By					Name of Program: TREATMENT OF HCV PATIENTS WITH DONATED "SOVALDI" MEDICINE IN THE REPUBLIC OF ARMENIA								
Print Name: TSOVINAR GALSTYAN													
Signature:					Name of Organisation: : "NORK" REPUBLICAN INFECTIOUS CLINICAL HOSPITAL								
Telephone Number:					Date aware of Safety Information: 26/06/2017								
Fax No/Email: galstyan.thsovinar@mail.ru					Country of Occurrence of Safety Information ARMENIA								
Patient Details													
DOB:         19/08/1982         (or year of birth):         Sex:         Male						e □ Female ☑ Initials: K.TS. Age: 34							
Drug Details (Provide additional drugs on a separate page)													
Lot/Batch No	In Reason For Taking Stop Date (or O			Τ	Start		Route		Dos	e	Drug Name		
TZDND	НЕР С	(DD/MON/YYYY)  On-going			(DD/MON/YYYY) NATURE (DD/MON/YYYYY) PO		РО	400MG		G G	SOVALDI		
Safety Information Details: Please provide a short summary of the adverse event(s) (AE) or other safety information (e.g. reports such as pregnancy, death, hospitalization, overdose, misuse, abuse, medication error, lack of effect, off-label use, occupational exposure, AEs associated with product complaints or AEs in an infant following exposure from breastfeeding). Please include the start and stop dates and the outcome of the event(s) or confirm if the event(s) is/are still ongoing. Please also provide any treatment given to treat the event(s), any relevant medical history and for reports of death include the date of death – continue on another page if necessary.													
Patient started treatment with Sovaldi on 01/06/2017. On 01.06.2017-02.06. 2017 he was experienced nausea, vomiting and general weakness. Events were recovered on 08.06.2017.													
After that patient mentioned weight decreased about 4 kg.													
Also patient mentioned about catarrhal events after the medicine intake.													
Has this safety information previously been reported to a Regulatory Authority? Yes □ No □					Does the Reporter consider that the event(s) were possibly related to the drug? Yes $\square$ No $\square$								
Reporter Details (i.e. who notified you of the above safety information?)													
Is the Reporter a: Doctor ☑ Nurse □ Pharmacist □ Non-healthcare professional (e.g. patient, relative)* □  If the Reporter is a Healthcare Professional (HCP) and they are willing to provide us with their contact information, please record below													
*If the Reporter is a Non-healthcare professional, please confirm if they are willing to provide contact information for their HCP:  Yes  (Please record HCP details below)  No													
HCP Address HCP Name:													
First Line:						HCP Telephone No/FAX No:							
Town/City:						·							
County/State:					HCP Email:								
Postcode/Zip code:													

Please be aware that information provided to the Ministry of Labour, Health and Social Affairs of Georgia(MoLHSA) relating to you, may be used to comply with applicable laws and regulations. By providing us with information you are consenting to the control and processing of this personal or sensitive data by MoLHSA in accordance with applicable data protection laws and the MoLHSA privacy policy.