District of Columbia DC Dual Choice Critical Incident Form

All incidents must be reported within 24 hours, or if received on weekend/holiday you may submit the next business day. Complete the form below with detailed information and attach to the **EnterpriseNow ticket.**

Member plan	MCO		Reason for report			
	UnitedHealthcare		☐ Adverse event			
UHC Dual Choice (D-SNP with LTSS)			☐ Serious reportable incident			
(B-ONF WIGHE 100)			☐ Reportable incident			
Member information						
Last name:		First name:		Date of birth:		
Gender: DO		DC Medicai	Medicaid ID #:			
UnitedHealthcare Medicaid ID #:				Waiver: □Yes □ No		
Incident information						
Occurrence of incident date:			Discovery of incident date:			
Time: (HH:MM a.m./p.m.)			Time: (HH:MM a.m./p.m.)			
Address of incident:			T			
City:			State:	ZIP code:		
Abuse Neglect or Explo	itation (ANE)					
Was ANE involved? ☐ Yes ☐ No			If yes, was incident reported? ☐ Yes ☐ No			
If yes, when was incident reported? Name of external agency ANE reported to:				ed to:		
Section 1 DC LTSS : Please select the appropriate adverse events category from the following list that most accurately describes the incident or event within a healthcare setting (waiver incidents listed in Section 2).						
Surgical or invasive procedure events						
☐ Surgery or other invasive procedure performed on the wrong site						
☐ Surgery or other invasive procedure performed on the wrong patient						
☐ Wrong surgical or other invasive procedure performed on a patient						
☐ Unintended retention of a foreign object in a patient after surgery or other invasive procedure						
☐ Intraoperative or immediately postoperative/postprocedural death in an ASA Class 1 patient						



Section 1 DC LTSS: Please select the appropriate **adverse events** category from the following list that most accurately describes the incident or event within a healthcare setting (waiver incidents listed in Section 2). **Product or device events** ☐ Patient death or serious injury associated with the use of contaminated drugs, devices or biologics provided by the healthcare setting ☐ Patient death or serious injury associated with the use of function of a device in patient care, in which the device is used or functions other than intended Patient death or serious injury associated with intravascular air embolism that occurs while being cared for in a healthcare setting **Patient protection events** ☐ Discharge or release of a patient/resident of any age, who is unable to make decisions, to other than an authorized person ☐ Patient death or serious injury associated with patient elopement (disappearance) ☐ Patient suicide, attempted suicide or self-harm that results in serious injury, while being cared for a health care setting **Care management events** ☐ Patient death or serious injury associated with a medication error (e.g., errors involving the wrong drug, wrong dose, wrong patient, wrong time, wrong rate, wrong preparation or wrong route of administration) ☐ Patient death or serious injury associated with unsafe administration of blood products ☐ Maternal death or serious injury associated with labor or delivery in a low-risk pregnancy while being cared for in a healthcare setting ☐ Death or serious injury of a neonate associated with labor or delivery in a low risk pregnancy ☐ Patient death or serious injury associated with a fall while being cared for in a healthcare setting Any Stage 3, Stage 4 and unstageable pressure ulcers acquired after admission/ presentation to a healthcare setting ☐ Artificial insemination with the wrong donor sperm or wrong egg ☐ Patient death or serious injury resulting from the irretrievable loss of an irreplaceable biological specimen ☐ Patient death or serious injury resulting from failure to follow up or communicate laboratory, pathology or radiology test results **Environmental event** ☐ Patient or staff death or serious injury associated with an electric shock in the course of a patient care process in a healthcare setting Any incident in which systems designated for oxygen or other gas to be delivered to a patient contains no gas, the wrong gas or are contaminated by toxic substances ☐ Patient or staff death or serious injury associated with a burn incurred from any source in the course of a patient care process in a healthcare setting ☐ Patient death or serious injury associated with the use of physical restraints or bedrails while being cared for in a healthcare setting



Section 1 DC LTSS (cont.) : Please select the appropriate adverse events category from the following list that most accurately describes the incident or event within a healthcare setting (waiver incidents listed in Section 2).				
Ra	diologic events			
	Death or serious injury of a patient or staff associated with the introduction of a metallic object into the MRI area			
Pot	tential criminal events			
	Any instance of care ordered by or provided by someone impersonating a physician, nurse, pharmacist or other licensed healthcare provider			
	Abduction of a patient/resident of any age			
	Sexual abuse/assault on a patient or staff member within or on the grounds of a health care setting			
	Death or serious injury of a patient or staff member resulting from a physical assault (i.e., battery) that occurs within or on the grounds of a healthcare setting			
Section 2 LTSS Waiver: Please select the appropriate reportable incident category from the following list that most accurately describes the incident or event regardless of location/setting.				
EPE	Waiver Serious Reportable Incidents (SRIs) SRIs include, but are not limited to:			
	Death			
	Abuse, neglect or exploitation			
	Theft of consumer personal property			
	Serious physical injury			
	Inappropriate or unauthorized use of restraints			
	Suicide attempt			
	Serious medication error			
	Hospitalization			
	Suicide threat			
	Vehicle accident			
	Fire or police involvement			
	Emergency room visit			
	Emergency relocation			
	Property destruction			
	Other events or situations that involve harm or risk of harm to a participant/member			



Section 2 LTSS Waiver (cont.): Please select the appropriate **reportable incident** category from the following list that most accurately describes the incident or event regardless of location/setting.

EPD Waiver Serious Reportable Incidents (SRIs) (cont.) SRIs include, but are not limited to:

Detailed description of incident:

Source of information					
Source of information for critical incident data					
Name:	Email:				
Phone number:					
Other individuals/witnesses					
Name:	Email:				
Phone number:					
Name:	Email:				
Phone number:					
Name:	Email:				
Phone number:					
External agencies contacted (APS, CPS, law enforcement, etc.)					
1. Agency:	Agency contact name:				
Phone number:	Date of report:				
2. Agency:	Agency contact name:				
Phone number:	Date of report:				
3. Agency:	Agency contact name:				
Phone number:	Date of report:				
Follow-up/resolution of incident					
Is the member subject to further harm, or do they have further emergency needs at this time? $\hfill \square$ Yes $\hfill \square$ No					



Detailed description of any/all follow-up action	s for this incident:			
Providers involved in incident				
Provider (1) name:	NPI number:			
Contact information:				
Address:				
City:	State:	ZIP code:		
Provider type:				
Provider (2) name:	NPI number:	NPI number:		
Contact information:				
Address:				
City:	State:	ZIP code:		

NPI number:

State:



ZIP code:

Provider type:

Provider type:

Address:

City:

Provider (3) name:

Contact information:

Follow-up/resolution of incident (cont.)

If yes, please explain