VCA - vascularized gland transplant recipient Uet TIED/* follow-up



OMB No. 0915-0157, Expiration date: 9/30/2026

The transplant recipient follow-up (TRF) forms are generated at six months, one year and annually thereafter following transplantation, until either graft failure, recipient death or lost to follow-up is reported. Forms are generated by the age at transplant, not the age at listing. The TRF record is completed by the transplant hospital responsible for follow-up of the recipient.

Complete the TRF with only the applicable patient information since the last follow-up period. It is not to contain information pertaining solely to the previous or next follow-up period. For example: the 6-month follow-up should contain information from the time after the TRR was completed to the 6-month transplant anniversary date; the 1-year follow-up should contain information from the day after the 6-month transplant anniversary date to the 1-year transplant anniversary date; the 2-year follow-up should contain information from the day after the 1-year transplant anniversary date to the 2-year anniversary date

Complete one TRF form for recipients of bilateral upper limbs. Complete separate TRF forms for each VCA organ transplant.

The TRF must be validated within 90 days of the record generation date. See OPTN Policies (https://optn.transplant.hrsa.gov/policies-bylaws/policies) for additional information.

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Question	Answers
TRF form type (prepopulated)*	
Recipient first name (prepopulated)*	
Recipient last name (prepopulated)*	
Recipient middle initial (prepopulated)	
Date of birth (prepopulated)*	
SSN (prepopulated)*	
Birth sex (prepopulated)*	
HIC (prepopulated)	
Transplant date (prepopulated)*	
State of permanent residence*	
Permanent ZIP code	
Expected date (prepopulated)*	

Provider information

Question	Answers
Treating reconstructive surgeon name*	<u> </u>
Treating reconstructive surgeon NPI #*	
Treating transplant physician name*	<u> </u>
Treating transplant physician NPI #*	
Follow-up care provided by*	
Other, specify	

Donor information

Question	Answers
UNOS donor ID # (prepopulated)*	

Donor type (prepopulated)*		
OPO (prepopulated)		
Patient status		
Question	Answers	
Date last seen, graft removed, or death*		
Patient status*		
If patient status is "Dead", select the patient's cause of death		
Primary cause of death		
Other, specify	-	
Has patient been hospitalized since the last patient status date*		
Number of hospitalizations (1-100)		
Socio-demographic information		
Question	Answers	
For recipients 18 years of age or older		
Working for income		
Casia damagnaphia information. Course of novement		
Socio-demographic information - Source of payment		
Question	Answers	
Grant funding*		
Institutional funding*		
Primary source of payment*		
If primary source of payment is "Foreign government, specify" select foreign government		
Primary source of payment - foreign government, specify		
Functional status		
Question	Answers	
For recipients younger than 18 years of age at transplant and younger than 26 years of age at follow-up		
Motor development		
Psychosocial consult performed*		

Functional status: SF-12 score - Physical health

Question	Answers
Physical functioning (PF) score (0.0-100.0)	
Role-physical (RP) score (0.0-100.0)	
Bodily pain (BP) score (0.0-100.0)	
General health (GH) score (0.0-100.0)	
Physical component summary (PCS) score (0.0-100.0)	

Functional status: SF-12 score - Mental health

Question	Answers
Vitality (VT) score (0.0-100.0)	
Social functioning (SF) score (0.0-100.0)	
Role-emotional (RE) score (0.0-100.0)	
Mental heath (MH) score (0.0-100.0)	
Mental component summary (MCS) score (0.0-100.0)	

Clinical information

Question	Answers
Enter height or height status	
Height (cm: 1.00-225.00)	·
Height status	
Enter weight or weight status	
Weight (kg: 0.45-294.84)*	
Weight status	

Clinical information: Noncompliance

Question	Answers
Immunosuppression*	
Rehabilitation*	
Level of activity*	
Other*	

Other, specify

Clinical information: Post-transplant

Question	Answers	
Graft status*		
If "Failed"		
Date of graft failure		
Causes of graft failure		
Acute rejection		
Acute rejection - Banff score		
Acute rejection - visual skin changes		
Chronic rejection		
Chronic rejection - visual skin changes		
Vascular complications		
Sepsis / infection		
Trauma		
Patient requested removal		•
Non-adherence		
Other		
Other, specify		
Did patient have any acute rejection episodes during the follow-up period		
If yes, number of episodes (1-100)		
Enter for each episode		
Date of acute rejection diagnosis		
Acute rejection was treated		
Visual skin changes		
Biopsy was done to confirm acute rejection		
Banff score		
Enter for each episode		
Date of acute rejection diagnosis		
Acute rejection was treated		
Visual skin changes		
Biopsy was done to confirm acute rejection		
Banff score		
Enter for each episode		
Date of acute rejection diagnosis		
Acute rejection was treated		
Visual skin changes		
Biopsy was done to confirm acute rejection		
Banff score		
Enter for each episode		
Date of acute rejection diagnosis		

		-
Visual skin changes		
Biopsy was done to confirm acute rejection		
Banff score		•
		•
Enter for each episode		
Date of acute rejection diagnosis		
Acute rejection was treated		
Visual skin changes		
Biopsy was done to confirm acute rejection		
Banff score		•
Clinical information: Most recent lab data	Answers	
	Allswers	
Enter serum creatinine or serum creatinine status Serum creatinine (mg/dL: 0.10-25.00)		
		•
Serum creatinine status		
Enter hemoglobin A1c or hemoglobin A1c status		
Hemoglobin A1c (%: 0-100)		-
Hemoglobin A1c status		•
Donor specific antibodies (DSA)*		
Clinical information: Complications	Answers	
	Answers	
Question	Answers	
Question New onset diabetes*	Answers	
Question New onset diabetes* Metabolic complications* Infectious complications*	Answers	-
Question New onset diabetes* Metabolic complications* Infectious complications* Other complications*	Answers	
Question New onset diabetes* Metabolic complications* Infectious complications*	Answers	
Question New onset diabetes* Metabolic complications* Infectious complications* Other complications*	Answers	
Question New onset diabetes* Metabolic complications* Infectious complications* Other complications*	Answers	
Question New onset diabetes* Metabolic complications* Infectious complications* Other complications*	Answers	· · · · · · · · · · · · · · · · · · ·
Question New onset diabetes* Metabolic complications* Infectious complications* Other complications*	Answers	
Question New onset diabetes* Metabolic complications* Infectious complications* Other complications*	Answers	· · · · · · · · · · · · · · · · · · ·
Question New onset diabetes* Metabolic complications* Infectious complications* Other complications*	Answers	
Question New onset diabetes* Metabolic complications* Infectious complications* Other complications*	Answers	
Question New onset diabetes* Metabolic complications* Infectious complications* Other complications*	Answers	
New onset diabetes* Metabolic complications* Infectious complications* Other complications* Other, specify Post-transplant malignancy		

Post-transplant malignancy: Donor related

Question	Answers
If post-transplant malignancy is yes	
Donor related	
If yes, select one or more tumor types and diagnosis date for each	
Skin: Squamous cell	Yes
Skin: Squamous cell - Diagnosis date	
Skin: Basal cell	Yes
Skin: Basal cell - Diagnosis date	
Skin: Melanoma	Yes
Skin: Melanoma - Diagnosis date	
Kaposi's sarcoma: Cutaneous	Yes
Kaposi's sarcoma: Cutaneous - Diagnosis date	
Kaposi's sarcoma: Visceral	Yes
Kaposi's sarcoma: Visceral - Diagnosis date	
Brain	
Brain - Other, specify	
Brain - Diagnosis date	
Renal carcinoma	Yes
Renal carcinoma - Diagnosis date	
Carcinoma of vulva, perineum or penis, scrotum	Yes
Carcinoma of vulva, perineum or penis, scrotum - Diagnosis date	
Carcinoma of the uterus	Yes
Carcinoma of the uterus - Diagnosis date	
Ovarian	Yes
Ovarian - Diagnosis date	
Testicular	Yes
Testicular - Diagnosis date	
Esophagus	Yes
Esophagus - Diagnosis date	
Stomach	Yes
Stomach - Diagnosis date	
Small intestine	Yes
Small intestine - Diagnosis date	
Pancreas	Yes

Pancreas - Diagnosis date	
Larynx	Yes
Larynx - Diagnosis date	
Tongue, throat	Yes
Tongue, throat - Diagnosis date	
Thyroid	Yes
Thyroid - Diagnosis date	
Bladder	Yes
Bladder - Diagnosis date	
Breast	Yes
Breast - Diagnosis date	
Prostate	Yes
Prostate - Diagnosis date	
Colo-rectal	Yes
Colo-rectal - Diagnosis date	
Primary hepatic tumor	Yes
Primary hepatic tumor - Diagnosis date	
Metastatic liver tumor	Yes
Metastatic liver tumor - Diagnosis date	
Lung	Yes
Lung - Diagnosis date	
Leukemia	Yes
Leukemia - Diagnosis date	
Sarcomas (excluding Kaposi's)	Yes
Sarcomas (excluding Kaposi's) - Diagnosis date	
Other cancers	Yes
Other cancers - Specify	
Other cancers - Diagnosis date	
Primary unknown	Yes
Primary unknown - Diagnosis date Post-transplant malignancy: Recurrence of pretransplant malignancy	
Question	Answers
If post-transplant malignancy is yes	
Recurrence of pre-transplant malignancy	
If yes, type of pre-existing tumor	

Type of pre-existing tumor		
If other cancer, specify		
Date of recurrence (post-transplant)		
Post-transplant malignancy: Post-transplant de novo solid tumor		
Question	Answers	
If post-transplant malignancy is yes		
De novo tumor		
If yes, select one or more tumor types and diagnosis date for each		
Skin: Squamous cell	Yes	
Skin: Squamous cell - Diagnosis date		
Skin: Basal cell		
	Yes	
Skin: Basal cell - Diagnosis date		
Skin: Melanoma	Yes	
Skin: Melanoma - Diagnosis date		
Kaposi's sarcoma: Cutaneous	Yes	
Kaposi's sarcoma: Cutaneous - Diagnosis date		
Kaposi's sarcoma: Visceral	Yes	
Kaposi's sarcoma: Visceral - Diagnosis date		
Brain		
Brain - Other, specify		
Brain - Diagnosis date		
Renal carcinoma	Yes	
Renal carcinoma - Diagnosis date	_	
Carcinoma of vulva, perineum or penis, scrotum	Yes	
Carcinoma of vulva, perineum or penis, scrotum - Diagnosis date		
Carcinoma of the uterus	Yes	
Carcinoma of the uterus - Diagnosis date		
Ovarian	Yes	
	L les	
Ovarian - Diagnosis date		
Testicular	Yes	
Testicular - Diagnosis date		
Esophagus	Yes	
Esophagus - Diagnosis date		
Stomach		
Comadi	Yes	
Stomach - Diagnosis date		

Small intestine	Yes
Small intestine - Diagnosis date	
Pancreas	Yes
Pancreas - Diagnosis date	
Larynx	Yes
Larynx - Diagnosis date	
Tongue, throat	Yes
Tongue, throat - Diagnosis date	
Thyroid	Yes
Thyroid - Diagnosis date	
Bladder	Yes
Bladder - Diagnosis date	
Breast	Yes
Breast - Diagnosis date	
Prostate	Yes
Prostate - Diagnosis date	
Colo-rectal Colo-rectal	Yes
Colo-rectal - Diagnosis date	
Primary hepatic tumor	Yes
Primary hepatic tumor - Diagnosis date	
Metastatic liver tumor	Yes
Metastatic liver tumor - Diagnosis date	
Lung	Yes
Lung - Diagnosis date	
Leukemia	Yes
Leukemia - Diagnosis date	
Sarcomas (excluding Kaposi's)	Yes
Sarcomas (excluding Kaposi's) - Diagnosis date	
Other cancers	Yes
Other cancers - Specify	
Other cancers - Diagnosis date	
Primary unknown	Yes
Primary unknown - Diagnosis date	

Post-transplant malignancy: Post-transplant lymphoproliferative disease and lymphoma

Question	Answers
If post-transplant malignancy is yes	
Post-transplant lymphoproliferative disease (PTLD) and Lymphoma	
Diagnosis date	
Pathology	
Other, specify	- <u></u>
Treatment	
Question	Answers
Antiviral*	
Antikiation	
Antibiotic*	
Antifungal*	
Topical immunosuppressive medication - Drugs use	d for acute rejection or maintenance
Question	Answers
Steroids (Clobetasol)	☐ Previous maintenance
	Current maintenance
	Anti-rejection indication
Tacrolimus (Protopic)	☐ Previous maintenance
	Current maintenance
	Anti-rejection indication
Other, specify 1	
Previous maintenance, current maintenance or anti-rejection	
1 Tevious maintenance, current maintenance or anti-rejection	☐ Previous maintenance ☐ Current maintenance
	Anti-rejection indication
Other, specify 2	
Previous maintenance, current maintenance or anti-rejection	Previous maintenance
	Current maintenance
	Anti-rejection indication
Non-topical immunosuppressive medication - Drug	used for induction, acute rejection, or maintenance
Question	Answers
Steroids (prednisone, methylprednisolone, Solumedrol, Medrol, Decadron)	Previous maintenance
	☐ Current maintenance ☐ Anti-rejection indication
	Anti-rejection indication

Question	Answers
Atgam	Anti-rejection indication
Campath (alemtuzumab, anti-CD52)	Anti-rejection indication
Cytoxan (cyclophosphamide)	Anti-rejection indication
Methotrexate (Folex PFS, Mexate-AQ, Rheumatrex)	☐ Anti-rejection indication
OKT3 (Orthoclone, muromonab)	Anti-rejection indication
Rituxan (rituximab)	☐ Anti-rejection indication
Simulect (basiliximab)	Anti-rejection indication
Thymoglobulin	☐ Anti-rejection indication
Non-topical immunosuppressive medication - Drug used for previo	ous maintenance or current maintenance
Cyclosporine, select from the following:	
FON (generic sudcensine)	

Question	Answers
Cyclosporine, select from the following:	
EON (generic cyclosporine)	☐ Previous maintenance ☐ Current maintenance
Gengraf (Abbott cyclosporine)	☐ Previous maintenance ☐ Current maintenance
Neoral (CyA-NOF)	Previous maintenance Current maintenance
Other generic cyclosporine, specify brand	
Other generic cyclosporine	☐ Previous maintenance ☐ Current maintenance
Sandimmune (cyclosporine A)	☐ Previous maintenance ☐ Current maintenance
Imuran (azathioprine, AZA)	☐ Previous maintenance ☐ Current maintenance
Leflunomide (LFL)	☐ Previous maintenance ☐ Current maintenance
Mycophenolate acid, select from the following:	
CellCept (MMF)	☐ Previous maintenance ☐ Current maintenance
Generic MMF (generic CellCept)	Previous maintenance Current maintenance

Myfortic (mycophenolate acid)	Previous maintenance
	Current maintenance
Nulojix (belatacept)	Previous maintenance
	Current maintenance
Rapamune (sirolimus, Rapamycin)	Previous maintenance
	Current maintenance
Tacrolimus, select from the following:	
Astagraf XL (extended release tacrolimus)	Previous maintenance
	Current maintenance
Generic tacrolimus (generic Prograf)	Previous maintenance
	Current maintenance
Prograf (FK506)	Previous maintenance
	Current maintenance
Zortress (everolimus)	Previous maintenance
	Current maintenance
Other drugs	
Other immunosuppressive medication 1, specify:	
Other immunosuppressive medication 1	Previous maintenance
	Current maintenance
	Anti-rejection indication
Other immunosuppressive medication 2, specify:	
Other immunosuppressive medication 2	Previous maintenance
	Current maintenance
	Anti-rejection indication

Public Burden/Privacy Act Statements

Department of Health and Human Services Health Resources and Services Administration OMB No: Expiration Date: 0915-0157 9/30/2026

ORGAN PROCUREMENT AND TRANSPLANTATION NETWORK (OPTN)

DATA COLLECTION

DATA ACCURACY CERTIFICATION: I certify that the data entered by me in UNetSM are accurate, timely, and complete to the best of my knowledge, information and belief. These data are based upon information contained in corresponding medical records and other source documents, or where appropriate, are based upon clinical observation.

PUBLIC BURDEN STATEMENT: An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. The OMB control number for this project is 0915-0157. Public reporting burden for the applicant for this collection of information is estimated to average 53 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to HRSA Reports Clearance Officer, 5600 Fishers Lane, Room 1033, Rockville, Maryland 20857.

PRIVACY ACT STATEMENT: In accordance with the requirements of the Privacy Act of 1974 (https://www.federalregister.gov/documents/2022/08/01/2022-16344/privacy-act-of-1974-system-of-records) as amended, 42 U.S.C. § 273, et seq., and 42 CRF Part 121 authorize collection of this information by the OPTN. This information is distributed to the Scientific Registry of Transplant Recipients (SRTR) and the Health Resources and Services Administration (HRSA), with the United States Department of Health and Human Services. The primary uses of this information are to match organ donors with recipients, to monitor compliance of member organizations with OPTN requirements, to review and report on the status of organ donation and transplantation in the United States, and to provide data to researchers and government agencies to study transplantation. The routine uses which may be made of this information are: (i) to organ procurement organizations and transplant hospitals to match organ donors with compatible recipients and validate the accuracy of donor and recipient; (ii) to the Department of Justice to use in defending litigation; (iii) to a congressional office upon the request of an individual concerning records pertaining to him/her; (iv) for research purposes, if certain requirements are satisfied and data use agreements are executed; and (v) to Agency contractors who have been engaged by the Agency to assist in accomplishment of an Agency function relating to the purposes of this system and who need to have access to the records in order to assist the Agency. Furnishing the remaining information requested is required by law of organ procurement organizations and transplant hospitals and the failure to submit such information may result in enforcement actions resulting from noncompliance with OPTN requirements. HRSA (08/02)

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