

## **Transfusion Recipient Database for the Recipient Epidemiology and Donor Evaluation Study-III (REDS-III)**

### **Documentation for Public Use Data files**

The objectives of the Recipient Epidemiology and Donor Evaluation Study-III (REDS-III) program are to ensure safe and effective blood banking and transfusion medicine practices through a comprehensive, multifaceted strategy involving basic, translational, and clinical research to improve the benefits of transfusion while reducing its risks. REDS-III was funded by the National Heart, Lung and Blood Institute of NIH.

To meet the objectives of REDS-III data were collected to produce three sets of core files. The donor/donation files contain information on attempted and successful donations of whole blood and blood products, as well as deferrals and the reasons for them. The component files contain information on processing of donated blood and blood products, such as separation of whole blood into red blood cells, platelets and plasma, as well as treatments, such as irradiation or leukocyte reduction. The transfusion recipient files contain information on both inpatient and outpatient transfusion recipients, such as type of product transfused, number of units transfused each day, and adverse outcomes for transfused patients. The three sets of files are being converted into three separate sets of public use files. One goal in REDS-III was to be able to link transfusion recipients to blood donations and blood processing records to permit studies of the effects of donor characteristics and component processing on transfusion outcomes. The ability to link the files has been preserved in the public use files. The remainder of this document is a discussion of the transfusion recipient files data and associated public use files.

Participants in the domestic component of the REDS-III program included:

- Four blood centers and 12 associated hospitals
  - Blood Center of Wisconsin
    - Froedtert Hospital, Milwaukee, WI
    - Marshfield Clinic
    - Saint Joseph's Ministry Hospital, Milwaukee, WI
    - Aurora Saint Luke's/Aurora Sinai Hospitals, Milwaukee, WI
  - Blood Centers of the Pacific
    - San Francisco General Hospital, San Francisco, CA
    - SF Veterans Administration Medical Center, San Francisco, CA
  - Institute for Transfusion Medicine
    - Presbyterian Hospital, Pittsburgh, PA
    - Shadyside Hospital, Pittsburgh, PA
    - St. Margaret's Hospital, Pittsburgh, PA
  - American Red Cross Blood Services - New England Region
    - Yale-New Haven Hospital, New Haven, CN
    - Bridgeport Hospital, Bridgeport, CN
- A data coordinating center (DCC)
  - RTI International; and
- A central laboratory
  - Blood Systems Research Institute in San Francisco

The REDS program began in 1989 primarily in response to the HIV epidemic. The focus of REDS and its successor, REDS-II was on studies of blood donors and blood donation. With the advent of REDS-III, a strong focus on transfusion recipients was added to studies of blood donors and donations. The new focus has included studies of blood component utilization, the effect of transfusion strategies on clinical outcomes in transfusion recipients and adverse effects

of transfusion over and above transfusion-transmitted infections. One important goal of REDS-III was to develop a database in which transfusion recipients could be linked to blood donors to examine the effects of donor characteristics on transfusion outcome. To this end, the REDS-III team collected data from 120,290 transfusion recipients over 234,277 encounters, where an encounter can be either an episode of outpatient transfusion or a hospitalization during which transfusion was received. Some recipients contributed data from a single encounter while others contributed data from two or more encounters. Multiple encounters with the same recipient may have been entirely outpatient visits, entirely hospitalizations or a mix of the two. Data were also collected from the transfusion recipients during hospitalizations in which they were not transfused. Data collection from recipients began in January 2013 and continued through December 2016. Additional data were collected during 1,285,359 encounters with nontransfused patients to provide a comparison group. Four subgroups of patients were defined during design of the data collection system:

- TX (Transfused): all eligible patients (inpatients and outpatients) that received a transfusion during an encounter.
- TS (NonTransfused-Type and Screened ONLY): all eligible patients (inpatients and outpatients) that underwent an antibody ID (ABID) and/or a type and screen ONLY during the encounter but did not receive a crossmatch or a transfusion.
- TCM (NonTransfused-Type-and-CrossMatched): all eligible patients (inpatients and outpatients) that underwent a type-and -crossmatch during the encounter but did not receive a transfusion.
- NR (a Non-Recipient who is non-transfused, not TS, not TCM): all eligible inpatients that did not receive a transfusion, nor a type-and-screen, nor a type-and-crossmatch during the inpatient encounter and did not have an eligible limited outpatient encounter in which a TS or TCM occurred within 45 days prior to admission.

#### The data collection process.

Shortly after REDS-III began, a working group of REDS-III investigators, DCC staff and representatives from NHLBI was assembled to develop the specifications for the data to be collected from blood donors. The original version of the data dictionary included 21 tables. Four tables were dropped for a variety of reasons, such as inability to scrub PHI at some hospitals and inability to report the data at others. One table was added. The final dataset included the 18 tables listed below.

1. RC\_BloodGas – Oxygen saturation, pH, CO<sub>2</sub> level, etc.
2. RC\_ChestStudy – outcome of CT, X-ray and MRI
3. RC\_CrossMatch – results of crossmatch, reported as compatible or incompatible
4. RC\_Demographic – date of birth, sex, race, ethnicity
5. RC\_DiagnosisCodes – ICD9 and ICD10 codes (both primary and secondary)
6. RC\_Diagnostics – results from a variety of diagnostic tests, such as urinalysis, CBC, albumin, or INR,
7. RC\_Encounters – encounter type (inpatient vs outpatient); for hospitalized patients, dates of admission and death or discharge with an indicator for outcome of the encounter, and length of ICU stay if placed in the ICU.
8. RC\_Fluids – intake or output of fluids, such as IV blood products, blood loss and urine output.

9. RC\_IssuedTXProducts – type of product issued for transfusion and patient location (emergency department, procedure suite, outpatient, etc.)
10. RC\_Medications – type of medication, dose and route of administration
11. RC\_MicrobialTests – type of specimen (blood, urine, etc.), type of culture (fungal, bacterial, other), results and identity of organisms if available
12. RC\_PreLabs – type of test (hemoglobin, UNI, platelet count) and measurement.
13. RC\_ProcedureCodes – ICD9 or ICD10
14. RC\_RespiratorySupport – ICD9 or ICD10 code for type of support, flow rate, FiO<sub>2</sub>.
15. RC\_Transfers – direction of transfer; location from which or to which patient was transferred
16. RC\_TxReactions – type of transfusion reaction (fever, edema, flush, hives, abdominal pain, etc.).
17. RC\_VentilatorDuration – length of time on ventilator support
18. RC\_Vitals – vital signs (blood pressure, arterial pressure, pulse, respiration, temperature, etc.)

Tables 8, RC\_Fluids, and 16, RC-TxReactions, are not included in the public use files because of problems with the reliability of the data at some sites. The other 16 tables are included in the public use files. Most of the tables included date and time, such as date and time of admission or discharge and date and time of blood draw. This is important information because some measurements or observations were recorded daily or repeatedly during a day in the hospital.

Data collection was more complete for transfused inpatients than for transfused outpatients or non-transfused inpatients. Data from NR patients was limited to blood gases if available, demographics, diagnosis codes, diagnostics, encounters, medications, pre-labs, procedure codes, respiratory support, transfers and vitals. A limited outpatient dataset (LOD) was defined to capture pre-admission testing of inpatients that occurred up to 45 days before an admission. The LOD included a subset of the variables from any outpatient visit where a type-and-screen, type-and-crossmatch and/or antibody ID testing occurred. The LOD included data from 7 tables: RC\_CrossMatch, RC\_Demographic, RC\_DiagnosisCodes, RC\_Diagnostics, RC\_Encounters, RC\_Medications, and RC\_ProcedureCodes. In addition, data from outpatient transfusions will be missing for procedures that are not routinely used in the outpatient setting.

The specifications for the data to be extracted from blood center records were sent to programmers at the four hubs. The programmers then developed extraction programs to pull the required data from the data management systems at the hubs. Data from the first quarter of 2013 were extracted for submission to the DCC. The data were reviewed at the hospitals before submission as an initial check against the specifications for data extraction provided by the DCC. The files were transmitted to the DCC through a secure portal that the DCC provided for this purpose. Further validation took place at the DCC, such as checks for duplicate records, range checks and value checks to make sure the values matched those allowed in the data dictionary. A report of the results of the checks was produced and sent to each hub for review. The hub programmers made any necessary changes to the extraction programs and sent revised data tables to the DCC for inspection and validation. Once this process was complete, data from the second quarter of 2013 were extracted and submitted to the DCC. The data were compared against data from the first quarter to ensure consistent application of the extraction rules.

Subsequent data collection involved extracting data using the same extraction program; reviewing data at the hub; uploading the data through the secure portal; reviewing data at the

DCC as described above, with queries back to the hub if needed; and loading the approved data into the donor database. The process of comparing data from a given quarter against data from previous quarters to ensure consistency of extraction continued throughout the data collection period.

Some of the files contain the results of diagnostic tests and other laboratory procedures. Rather than a separate field for each test, the files include one record for each test that was performed, with test type, resulting value and unit listed in separate fields. Separate records are provided if the same test was performed two or more times on a subject. If some tests produce quantitative results and others produce qualitative results, then the two types of results are in separate fields. For some tests, the format of the unit of measure varied among sites, such as g/dl vs. g/dL or G/DL. The units themselves also varied among sites, such as height in inches vs cm or concentration in mmol vs grams. Many of these problems were eliminated during production of the public use files. However, the user is cautioned to review the data before proceeding with an analysis.

#### The public use files

Some recipient files contained information obtained once during an encounter while others contained information collected more than once, possibly daily, or even several times a day. A new record was generated each time data were collected, which means that the number of records per encounter varied quite widely recipients and files. Because of this variation in structure, the files were, for the most part, kept separate when the public use versions were produced. Only RC\_Demographic and RC\_Encounters were merged into a single file. The vital signs table is so large that it was split into two files, each with half the total observations.

#### Changes to the data

The process of anonymizing the files involved replacing some variables and deleting others, as described below. It also proved necessary to add some variables to preserve the longitudinal structure of the data.

The recipient data files included several identifiers: subjectID, hospitalcode, blood center, encounterID, donation identification number (DIN) and productkey. The first five were replaced with random numbers. The replacement variables are SubjectID Random, HospitalCode Random, CenterID Random, EncounterID Random, and DIN Random respectively. To preserve the ability to link records and cluster encounters within subject, all occurrences of a given value for one of the identifiers were replaced with the same random number. To preserve the ability to link donation records to transfusion recipients, the random replacement for the DIN in the donor files was also used to replace the same DIN in the recipient files. Productkey contained both the DIN and the product code. The two components were separated into two variables. The DIN in the product key was replaced with DIN PK Random, which is included in REDS\_III\_RC\_ISSUEDTXPRODUCTS\_PUD.SAS7BDAT.

The user should be aware that the blood center that supplied a product to the hospital may not be the point of origin of that product. Some of the blood centers involved in REDS-III imported some blood products from other domestic blood centers. Imported products are identified in the imports table of the components database.

In the original files, all dates were stored as three separate fields - month, day, and year. Date of birth was replaced with integer age at the time of contact. For all other dates, day was deleted but month and year were retained in the files. Two variables were added to the files to preserve the longitudinal structure of the data. Days since the start of a given encounter was recorded as DaysSinceStartEncounter. This variable takes the value 0 on the day the encounter starts, 1 the next day and so on. DaysSinceStart1stEncounter represents days from the start of the first encounter for a recipient to the start of each subsequent encounter for the same recipient. All records in an encounter were assigned the same value. For example, if a second encounter started 14 days after the start of the first encounter and lasted 6 days with daily data collection, then DaysSinceStart1stEncounter took the value 14 for all records in the second encounter. Time since the start of the first encounter to any day within a second or subsequent encounter is the sum of DaysSinceStartEncounter and DaysSinceStart1stEncounter. The duration of an encounter is recorded as DaysToEncounterEndDate.

Many of the observations in the recipient database included the time as well as month, day and year. Examples include timing of blood draw, time of initiation of ventilator support and time of administration of a medication. In many cases, the same event occurred more than once on a given day. Time of day was preserved in the public use files to allow the user to properly order the events.

Storage duration prior to transfusion is an issue of potential interest in some analyses. Therefore, DaysDonationToIssue was added to REDS\_III\_RC\_ISSUEDTXPRODUCTS\_PUD.SAS7BDAT to represent days between donation and product issue for transfusion. This was calculated using date of donation from the REDS-III donation database and date of product issue from the issued products table of the recipient database.

In some cases, a transfusion involved pooled blood products, such as pooled platelets. A binary indicator, DINPooledLinkFlag, was added to the issued products table to identify these units. Further information on pooled products can be obtained from the components database.

Sixteen public use files were produced after these changes were implemented:

1. REDS\_III\_RC\_BloodGas\_PUD.SAS7BDAT – Oxygen saturation, pH, CO<sub>2</sub> level, etc.
2. REDS\_III\_RC\_ChestStudy\_PUD.SAS7BDAT – outcome of CT, X-ray and MRI
3. REDS\_III\_RC\_CrossMatch\_PUD.SAS7BDAT – results of crossmatch, reported as compatible or incompatible
4. REDS\_III\_RC\_DiagnosisCodes\_PUD.SAS7BDAT – ICD9 and ICD10 codes (both primary and secondary)
5. REDS\_III\_RC\_Diagnostics\_PUD.SAS7BDAT – results from a variety of diagnostic tests, such as urinalysis, CBC, albumin, or INR,
6. REDS\_III\_RC\_Encounterdemographic.SAS7BDAT – encounter type (inpatient vs outpatient); for hospitalized patients, dates of admission and death or discharge with an indicator for outcome of the encounter, and length of ICU stay if placed in the ICU, etc. Also, – date of birth, sex, race, ethnicity, etc
7. REDS\_III\_RC\_IssuedTXProducts\_PUD.SAS7BDAT – type of product issued for transfusion and patient location (emergency department, procedure suite, outpatient, etc.)

8. REDS\_III\_RC\_Medications\_PUD.SAS7BDAT – type of medication, dose and route of administration
9. REDS\_III\_RC\_MicrobialTests\_PUD.SAS7BDAT – type of specimen (blood, urine, etc.), type of culture (fungal, bacterial, other), results and identity of organisms if available
10. REDS\_III\_RC\_PreLabs\_PUD.SAS7BDAT – type of test (hemoglobin, UNI, platelet count) and measurement.
11. REDS\_III\_RC\_ProcedureCodes\_PUD.SAS7BDAT – ICD9 or ICD10
12. REDS\_III\_RC\_RespiratorySupport.SAS7BDAT – ICD9 or ICD10 code for type of support, flow rate, FiO<sub>2</sub>.
13. REDS\_III\_RC\_Transfers\_PUD.SAS7BDAT – direction of transfer; location from which or to which patient was transferred
14. REDS\_III\_RC\_VentilatorDuration.SAS7BDAT – length of time on ventilator support
15. RC\_Vitals\_PUD\_1.SAS7BDAT and RC\_Vitals\_PUD\_2.SAS7BDAT – vital signs (blood pressure, arterial pressure, pulse, respiration, temperature, etc.)

Further details on content are provided in the codebook and detailed data dictionary.

#### Merging the files

If two files contain one record per encounter, then they can be merged using SubjectID\_Random and EncounterID\_random. Additional information may be required if one or more files contain multiple records from the same encounter. In that case, the user should consider adding DaysSinceStartEncounter and time to the list of variables used to merge records.

The replacements for identifying variables were constructed to preserve the ability to link records in the recipient database to records in the donor/donation and components databases. Using the variables for this purpose is covered in another document.

#### Problems with the data

While extensive effort was made to collect complete data, some data are missing from the files. This does not include data missing by design; i.e. data on forms not used on some subgroups of subjects, as described earlier.

Users should also be aware that some fields contain small proportions of anomalous values. Examples include hemoglobin levels that are outside the range expected for blood donors and age at donation that is either too young for the donor to be eligible to donate or too old to be realistic. The problems are mainly the result of data entry errors at the participating hospitals or blood centers; i.e. the data were downloaded as is from the hospitals and blood center files. Users are strongly encouraged to run univariate statistics on all variable of interest to identify values that are likely erroneous before proceeding with data analysis.

## **APPENDIX**

**Specifications for the data collection tables for the REDS-III transfusion recipient database**

Specifications for RC\_BloodGas

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
EncounterID		ID for Encounter. Will appear on all records in all tables for a specific encounter.	Int	10		R		Y	Y	Y	Y
SubjectID		SubjectID will uniquely identify subject and will appear on all records in all tables.	Int	13		R		Y		Y	Y
DateD		Date of draw - Day	Int	2	dd	R		Y	Y	Y	Y
	1-31										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
DateM		Date of draw - Month	Int	2	mm	R		Y	Y	Y	Y
	1-12										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
DateY		Date of draw - Year	Int	4	yyyy	R		Y	Y	Y	Y
	2012-2020										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
Time		Time of draw	Time	5	hh:mm	R		Y	Y	Y	Y
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
SAO2	ABG: 75 - 100 VBG: 20 - 100	Value - SAO2 / SvO2	Int	3	%			Y	Y	Y	Y
pH	6.7 – 7.7	Value - pH	Single	3.2				Y	Y	Y	Y
PCO2	20-100	Value - PCO2	Int	3	mmHg			Y	Y	Y	Y

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
PO2	ABG: 40 - 500 VBG: 20 - 100	Value - PO2 / PvO2	Int	3	mmHg			Y	Y	Y	Y
HCO3	5-40	Value - HCO3	Int	2	mmol/L			Y	Y	Y	Y
ABE	-15 to 15	Value - ABE	Single	(+/-) 3.1	mmol/L			Y	Y	Y	Y
BGType		Type of Blood Gas (Arterial/Venous)	Int	2		R		Y	Y	Y	Y
	1	Arterial									
	2	Venous									
FlowRate	0.5 – 15.0	FlowRate	Single	3.1	L/min			Y	Y	Y	Y
FiO2	21 - 100	Value - FiO2	Int	3	%			Y	Y	Y	Y

Specifications for RC\_ChestStudy

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
EncounterID		ID for Encounter. Will appear on all records in all tables for a specific encounter.	Int	10		R			Y	Y	Y
SubjectID		SubjectID will uniquely identify subject and will appear on all records in all tables.	Int	13		R			Y	Y	Y
ChestStudyType		Type of radiographic chest study	Int	2		R			Y	Y	Y
	1	Chest X-Ray									
	2	Chest CT									
	3	Chest MRI									
Indication			Char	1500					Y	Y	Y
Interpretation		Leave blank if PHI is present.	Char	50000					Y	Y	Y
DateD		Date radiographic chest study performed- Day	Int	2	dd	R			Y	Y	Y
	1-31										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
DateM		Date radiographic chest study performed - Month	Int	2	mm	R			Y	Y	Y
	1-12										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
DateY		Date radiographic chest study performed - Year	Int	4	yyyy	R			Y	Y	Y
	2012-2020										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
Time		Time radiographic chest study performed	Time	5	hh:mm	R			Y	Y	Y



Specifications for RC\_CrossMatch

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
EncounterID		ID for Encounter. Will appear on all records in all tables for a specific encounter.	Int	10		R	Y			Y	Y
SubjectID		SubjectID will uniquely identify subject and will appear on all records in all tables.	Int	13		R	Y			Y	Y
ProductKey		ID to uniquely identify Product crossmatched. <b>(For ISBT and Codabar: DIN + Product Code)</b>	Char	21		R	Y			Y	Y
CMDateD		Crossmatch Date - Day	Int	2	dd	R	Y			Y	Y
	1-31										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
CMDateM		Crossmatch Date - Month	Int	2	mm	R	Y			Y	Y
	1-12										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
CMDateY		Crossmatch Date - Year	Int	4	yyyy	R	Y			Y	Y
	2012-2020										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
CMTIME		Crossmatch Time	Time	5	hh:mm	R	Y			Y	Y
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									



## Specifications for RC\_Demographic



## Specifications for RC\_DiagnosisCodes



## Specifications for RC\_Diagnostics

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
EncounterID		ID for Encounter. Will appear on all records in all tables for a specific encounter.	Int	10		R	Y	Y	Y	Y	Y
SubjectID		SubjectID will uniquely identify subject and will appear on all records in all tables.	Int	13		R	Y	Y	Y	Y	Y
DiagnosticType		Type of diagnostic lab test	Int	2		R	Y	Y	Y	Y	Y
	1	Patient ABO					Y	Y	Y	Y	Y
	2	Patient Rh					Y	Y	Y	Y	Y
	3	AbID-Screen					Y		Y	Y	Y
	4	AbID-RBC					Y		Y	Y	Y
	5	AbID-Eluate					Y		Y	Y	Y
	6	DATPolyspecific					Y		Y	Y	Y
	7	DATIGG					Y		Y	Y	Y
	8	DATComplement					Y		Y	Y	Y
	9	INR					Y	Y	Y	Y	Y
	10	aPTT							Y	Y	Y
	11	Hgb					Y	Y	Y	Y	Y
	12	Fibrinogen					Y	Y	Y	Y	Y
	13	White Blood Cells (WBC)					Y	Y	Y	Y	Y
	14	Platelet count					Y	Y	Y	Y	Y
	15	Creatinine					Y	Y	Y	Y	Y
	16	Total bilirubin					Y	Y	Y	Y	Y
	17	Direct bilirubin					Y	Y	Y	Y	Y
	18	Haptoglobin							Y	Y	Y
	19	LDH							Y	Y	Y

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
	20	ALT							Y	Y	Y
	21	AST							Y	Y	Y
	22	Albumin					Y	Y	Y	Y	Y
	23	Lactate					Y	Y	Y	Y	Y
	24	BNP						Y	Y	Y	Y
	25	pro-BNP						Y	Y	Y	Y
	26	Sodium					Y	Y	Y	Y	Y
	27	Blood Urea Nitrogen (BUN)					Y	Y	Y	Y	Y
	28	Serum Glucose					Y	Y	Y	Y	Y
	29	Hematocrit					Y	Y	Y	Y	Y
	30	Troponin I					Y	Y	Y	Y	Y
	31	d-dimers						Y	Y	Y	Y
	32	Urinalysis - Glucose						Y	Y	Y	Y
	33	Urinalysis - Ketones						Y	Y	Y	Y
	34	Urinalysis - Occult Blood						Y	Y	Y	Y
	35	Urinalysis - Protein						Y	Y	Y	Y
	36	Urinalysis - Nitrite						Y	Y	Y	Y
	37	Urinalysis - Bilirubin						Y	Y	Y	Y
	38	Urinalysis - Specific Gravity						Y	Y	Y	Y
	39	Urinalysis - pH Urine						Y	Y	Y	Y
	40	Urinalysis - Urobilinogen						Y	Y	Y	Y
	41	Urinalysis - Urine Leukocyte Esterase						Y	Y	Y	Y
	42	Urinalysis - Squamous Epithelial Cells						Y	Y	Y	Y
	43	Urinalysis - Leukocyte						Y	Y	Y	Y
	44	Urinalysis - Hyaline Casts						Y	Y	Y	Y
	45	Urinalysis - Erythrocytes						Y	Y	Y	Y
	46	Urinalysis - Bacteria						Y	Y	Y	Y
	47	Activated Protein C Resistance						Y	Y	Y	Y

FieldName	Valid Values	Description	Data Type	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
	48	Factor V Leiden							Y	Y	Y
	49	Prothrombin Gene G20210A Mutation							Y	Y	Y
	50	Protein C Activity - amount							Y	Y	Y
	51	Protein C Activity - percent							Y	Y	Y
	52	Protein C Antigen - amount							Y	Y	Y
	53	Protein C Antigen - percent							Y	Y	Y
	54	Protein S Activity - amount							Y	Y	Y
	55	Protein S Activity - percent							Y	Y	Y
	56	Protein S Antigen (total) - amount							Y	Y	Y
	57	Protein S Antigen (total) - percent							Y	Y	Y
	58	Protein S Antigen (free) - amount							Y	Y	Y
	59	Protein S Antigen (free) - percent							Y	Y	Y
	60	Antithrombin III Activity - amount							Y	Y	Y
	61	Antithrombin III Activity - percent							Y	Y	Y
	62	Antithrombin III Antigen - amount							Y	Y	Y
	63	Antithrombin III Antigen - percent							Y	Y	Y
	64	Lupus Anticoagulant-DRVVT or DRVV (Quantitative)							Y	Y	Y
	65	Lupus Anticoagulant-DRVVT Abnormal Flag (Qualitative)							Y	Y	Y
	66	Lupus Anticoagulant HPPN (Quantitative)							Y	Y	Y
	67	Lupus Anticoagulant HPPN Abnormal Flag (Qualitative)							Y	Y	Y
	68	Homocysteine							Y	Y	Y
	69	Thrombin Time							Y	Y	Y
	70	Cardiolipin antibodies - IgA							Y	Y	Y
	71	Cardiolipin antibodies - IgG							Y	Y	Y
	72	Cardiolipin antibodies - IgM							Y	Y	Y
	73	Beta 2 Microglobulin							Y	Y	Y



FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
	-9	NOT COLLECTED at Hospital or Hub									
DrawDateD		Date of Draw (Collection) - Day	Int	2	dd	R	Y	Y	Y	Y	Y
	1-31										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
DrawDateM		Date of Draw (Collection) - Month	Int	2	mm	R	Y	Y	Y	Y	Y
	1-12										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
DrawDateY		Date of Draw (Collection) - Year	Int	4	yyyy	R	Y	Y	Y	Y	Y
	2012-2020										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
DrawTime		Time of Draw (Collection)	Time	5	hh:mm	R	Y	Y	Y	Y	Y
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
LabValue		Quantitative value of lab test results	Double	9.3				Y	Y	Y	Y
LabUnit		Unit of measure	Char	9				Y	Y	Y	Y



FieldName	Valid Values	Description	Data-Type	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
	NEG										
	POS										
	TRACE										
	NON-SPECIFIC										
	1+										
	2+										
	3+										
	4+										
	O										
	A										
	B										
	AB										
	UNT										
	<i>antibody name from attached table of antibodies</i>							Y	Y	Y	
	-6	Other Notation						-	-	-	-

## Specifications for RC Encounters





FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
	-9	NOT COLLECTED at Hospital or Hub									
DischargeTime		<b>Discharge Time</b>	Time	5	hh:mm	R	Y	Y	Y	Y	Y
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
VentilatorNeeded		<b>Derived: Invasive mechanical ventilation (y/n)</b>	Char	2			Y	Y	Y	Y	Y
	Y	Yes									
	N	No									
VentilatorDays		<b>Calculated: days of mechanical ventilation support</b>	Single	4.1	days				Y	Y	Y
VentilatorFreeDays		<b>Calculated: 28 - days of mechanical ventilation support</b>	Single	4.1	days				Y	Y	Y
ICU_LOS		<b>Calculated: Total amount of time the patient was in the ICU during the encounter (days).</b>	Single	4.1	days				Y	Y	Y
ICUFreeDays		<b>Calculated: 28 days - ICU LOS</b>	Single	4.1	days				Y	Y	Y
Mortality		<b>Hospital mortality at discharge</b>	Int	2		R	Y	Y	Y	Y	Y
	1	Alive									
	2	Death									
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
DeathDateD		<b>Death Date - Day</b>	Int	2	dd				Y	Y	Y

FieldName	Valid Values	Description	Data Type	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
	1-31										
<b>DeathDateM</b>		<b>Death Date - Month</b>	Int	2	mm			Y	Y	Y	Y
	1-12										
<b>DeathDateY</b>		<b>Death Date - Year</b>	Int	4	yyyy			Y	Y	Y	Y
	2012-2020										

Specifications for RC\_Fluids

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
EncounterID		ID for Encounter. Will appear on all records in all tables for a specific encounter.	Int	10		R			Y	Y	Y
SubjectID		SubjectID will uniquely identify subject and will appear on all records in all tables.	Int	13		R			Y	Y	Y
FluidType		Defines type of fluid measurement	Int	2		R		Y	Y	Y	
	1	Intake - Enteral									
	2	Intake - Intravenous: Crystalloids									
	3	Intake - Intravenous: Colloids									
	4	Intake - Intravenous: Blood products									
	5	Intake - Intravenous: Nutritional products									
	6	Intake - Intravenous: All medication infusions									
	7	Intake - Other									
	8	Intake - OR Total Fluid Intake									
	9	Output - Blood									
	10	Output - Urine									
	11	Output - Renal replacement therapy									
	12	Output - Other									
	13	Output - OR Total Fluid Out									
Value		Value - intake/output volume	Int	5	mL	R		Y	Y	Y	
	0-10000										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
DateD		Date of fluid measurement - Day	Int	2	dd	R			Y	Y	Y





Specifications for RC\_IssuedTXProducts

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
EncounterID		ID for Encounter. Will appear on all records in all tables for a specific encounter.	Int	10		R					Y
SubjectID		SubjectID will uniquely identify subject and will appear on all records in all tables.	Int	13		R					Y
ProductKey		ID to uniquely identify each Product in table. ( For ISBT and Codabar: DIN + Product Code)	Char	21		R					Y
IssueDateD		Date product issued to patient - Day	Int	2	dd	R					Y
	1-31										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
IssueDateM		Date product issued to patient - Month	Int	2	mm	R					Y
	1-12										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
IssueDateY		Date product issued to patient - Year	Int	4	yyyy	R					Y
	2012-2020										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
IssueTime		Time product issued to patient	Time	5	hh:mm	R					Y



## Specifications for RC\_Medications

FieldName	Valid Values	Description	Data Type	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
MedAdminDateY		<b>Date of medication administration - Year</b>	Int	4	yyyy	R	Y	Y	Y	Y	Y
	2012-2020										
	-3	Not Applicable									
	-8	NOT REPORTED or UNKNOWN for Participant									
MedAdminTime		<b>Time of medication administration</b>	Time	5	hh:mm	R	Y	Y	Y	Y	Y
	-3	Not Applicable									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
MedStopDateD		<b>Stop Date of medication administration - Day</b>	Int	2	dd	R	Y	Y	Y	Y	Y
	1-31										
	-3	Not Applicable									
	-8	NOT REPORTED or UNKNOWN for Participant									
MedStopDateM		<b>Stop Date of medication administration - Month</b>	Int	2	mm	R	Y	Y	Y	Y	Y
	1-12										
	-3	Not Applicable									
	-8	NOT REPORTED or UNKNOWN for Participant									
MedStopDateY		<b>Stop Date of medication administration - Year</b>	Int	4	yyyy	R	Y	Y	Y	Y	Y
	2012-2020										
	-3	Not Applicable									
	-8	NOT REPORTED or UNKNOWN for Participant									
MedStopTime		<b>Stop Time of medication administration</b>	Time	5	hh:mm	R	Y	Y	Y	Y	Y









## Specifications for RC\_MicrobialTests

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
	-9	NOT COLLECTED at Hospital or Hub									
CultureSource		Specimen source	Int	2		R		Y	Y	Y	Y
	1	Blood									
	2	Urine									
	3	Wound									
	4	CSF (cerebrospinal fluid)									
	5	Sputum									
	6	Bronchoalveolar lavage									
	7	Sinus									
	8	Tissue/biopsy									
	9	Stool									
	10	Other									
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
CultureResults		Culture Results	Int	2		R		Y	Y	Y	Y
	1	Growth									
	2	No Growth									
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
Organism1		Record up to six organisms; see attached table of organism codes	Int	3					Y	Y	Y
Organism2			Int	3					Y	Y	Y
Organism3			Int	3					Y	Y	Y
Organism4			Int	3					Y	Y	Y

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
Organism5			Int	3					Y	Y	Y
Organism6			Int	3					Y	Y	Y

Specifications for RC\_PreLabs

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
SubjectID		SubjectID will uniquely identify subject and will appear on all records in all tables.	Int	13		R		Y	Y	Y	Y
DiagnosticType		Type of diagnostic lab test	Int	2		R		Y	Y	Y	Y
	9	INR						Y	Y	Y	Y
	11	Hgb						Y	Y	Y	Y
	14	Platelet count						Y	Y	Y	Y
DrawDateD		Date of Draw (Collection) - Day	Int	2	dd	R		Y	Y	Y	Y
	1-31										
DrawDateM		Date of Draw (Collection) - Month	Int	2	mm	R		Y	Y	Y	Y
	1-12										
DrawDateY		Date of Draw (Collection) - Year	Int	4	yyyy	R		Y	Y	Y	Y
	2012-2020										
DrawTime		Time of Draw (Collection)	Time	5	hh:mm	R		Y	Y	Y	Y
LabValue		Quantitative value of lab test results	Double	9.3		R		Y	Y	Y	Y
LabUnit		Unit of measure	Char	9				Y	Y	Y	Y
	sec										
	g/dL										
	mg/dL										
	K/uL										
	U/L										
	mmol/L										
	pg/mL										
	Meq/L										
	/HPF										
	mg/L FEU										
	ug/mL DDU										



## Specifications for RC\_ProcedureCodes



## Specifications for RC\_RespiratorySupport

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
	4	Double Flow, Downs' Flow, and Optiflow									
	5	CPAP									
	6	BiPAP									
	7	Invasive mechanical									
	8	Room air									
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
FlowRate	0.5 – 15.0	Flow Rate							Y	Y	Y
	-6	Other Notation							Y	Y	Y
	-8	NOT REPORTED or UNKNOWN for Participant							Y	Y	Y
	-9	NOT COLLECTED at Hospital or Hub							Y	Y	Y
FiO2	21-100	FiO2 value							Y	Y	Y
	-6	Other Notation							Y	Y	Y
	-8	NOT REPORTED or UNKNOWN for Participant							Y	Y	Y
	-9	NOT COLLECTED at Hospital or Hub							Y	Y	Y

## Specifications for RC\_Transfers

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
	60	Outpatient									
	70	Recovery Room									
	80	Home									
	90	Nursing Home									
	100	Another Hospital									
	110	Long Term Assisted Care (LTAC)									
	120	Hospice									
	130	Rehab									
	190	Unknown									
	200	Other (not one of the defined locations)									
DateD		Date of transfer - Day							Y	Y	Y
	1-31										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
DateM		Date of transfer - Month							Y	Y	Y
	1-12										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
DateY		Date of transfer - Year							Y	Y	Y
	2012-2020										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
Time		Time of transfer							Y	Y	Y
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									

## Specifications for RC\_TxReactions

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
DOB		Birthdate of subject	Date	10	mm/dd/yyyy	R					Y
	01-12	Month									
	01-31	Day									
	1900-2012	Year									
x fusReason		Primary Reason for Transfusion	Char	9		R					Y
	COAG	Coagulopathy									
	GENDISORD	Genetic Disorder									
	HEMDISORD	Hematology Disorder									
	HEMOLYSIS	Hemolysis									
	INTBLEED	Internal Bleeding									
	MALIG	Malignancy									
	MEDICAL	Medical									
	SURGERY	Surgery									
	UNKNOWN	Unknown									
	OTHER	Other reason for transfusion									
x fusReasonSfy		Primary Reason for Transfusion - Specify Other	Char	100							Y
AdvRxnDate		Date of reaction	Date	10	mm/dd/yyyy	R					Y
	01-12	Date of reaction - Month									
	01-31	Date of reaction - Day									
	2012-2020	Date of reaction - Year									
AdvRxnTime		Time of reaction	Time	5		R					Y
Advrxtimetimeunk	Y/N	Time of reaction - Unknown	Char	1							Y
ss_biprlow	Y/N	Blood pressure decrease	Char	1							Y
ss_shock	Y/N	Shock	Char	1							Y
ss_coag	Y/N	Disseminated intravascular coagulation	Char	1							Y

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
ss_hemoglob	Y/N	Hemoglobinemia	Char	1							Y
ss_screen	Y/N	Positive antibody screen	Char	1							Y
ss_chills	Y/N	Chills/rigor	Char	1							Y
ss_fever	Y/N	Fever	Char	1							Y
ss_edema	Y/N	Edema	Char	1							Y
ss_flush	Y/N	Flushing	Char	1							Y
ss_jaundc	Y/N	Jaundice	Char	1							Y
ss_othrashh	Y/N	Other rash	Char	1							Y
ss_itch	Y/N	Pruritis (itching)	Char	1							Y
ss_hives	Y/N	Urticaria (hives)	Char	1							Y
ss_hematuri	Y/N	Hematuria	Char	1							Y
ss_hemognur	Y/N	Hemoglobinuria	Char	1							Y
ss_oliguria	Y/N	Oliguria	Char	1							Y
ss_abdpain	Y/N	Abdominal pain	Char	1							Y
ss_backpain	Y/N	Back pain	Char	1							Y
ss_flnkpain	Y/N	Flank pain	Char	1							Y
ss_sitepain	Y/N	Infusion site pain	Char	1							Y
ss_infiltr	Y/N	Bilateral infiltrates on chest x-ray	Char	1							Y
ss_broncspas	Y/N	Bronchospasm	Char	1							Y
ss_cough	Y/N	Cough	Char	1							Y
ss_hypox	Y/N	Hypoxemia	Char	1							Y
ss_shortBr	Y/N	Shortness of breath	Char	1							Y
ss_othersn	Y/N	Other signs/symptoms	Char	1							Y
otherSignSfy		Other signs/symptoms, specify	Char	100							Y
advRxn		Adverse Reaction	Char	6		R					Y

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
	ALLERG	Allergic reaction, including anaphylaxis									
	AHTR	Acute hemolytic transfusion reaction (AHTR)									
	DHTR	Delayed hemolytic transfusion reaction (DHTR)									
	DSTR	Delayed serologic transfusion reaction (DSTR)									
	FNHTR	Febrile non-hemolytic transfusion reaction(FNHTR)									
	HTR	Hypotensive transfusion reaction									
	INF	Infection									
	PTP	Post tranfusion purpura									
	TACO	Transfusion associated circulatory overload (TACO)									
	TAD	Transfusion associated dyspnea (TADO)									
	TA-GVND	Transfusion aossociated graft vs. host disease (TA_GVHD)									
	TRALI	Transfusion related acute lung injury (TRALI) antibody studies									
	UNK	Unknown pathophysiology									
	OTHER	Other Adverse Reaction									
OtherAdvRxn		Other Adverse Reaction, specify	Char	100							Y
ImAHTR	Y/N	AHTR Immune	Char	1							Y
ImmuneABAHTR		AHTR Antibody	Char	12							Y
NonimAHTR	Y/N	AHTR Non-Immune	Char	1							Y
NonimAHTRSfy		AHTR Non-Immune, Specify	Char	50							Y
ImDHTR	Y/N	DHTR Immune	Char	1							Y
ImmuneABDHTR		DHTR Antibody	Char	12							Y

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
NonimDHTR	Y/N	DHTR Non-Immune	Char	1							Y
NonimDHTRSfy		DHTR Non-Immune, Specify	Char	50							Y
ImmuneABDSTR		DSTR Antibody	Char	12							Y
rcpntTestPerform	Y/N	Infection:Test to detect specific pathogen on recipient post-Tx	Char	1							Y
rcpntTestPositive	Y/N	Infection:Recipient - Positive/Negative results	Char	1							Y
rcpntPathDesc1		Infection:Recipient Org1	Char	7							Y
rcpntPathDesc2		Infection:Recipient Org2	Char	7							Y
rcpntPathDesc3		Infection:Recipient Org3	Char	7							Y
donorTestPerform	Y/N	Infection:Test to detect specific pathogen on donor post-Tx	Char	1							Y
donorTestPositive	Y/N	Infection:Donor - Positive/Negative results	Char	1							Y
donorPathDesc1		Infection:Donor Org1	Char	7							Y
donorPathDesc2		Infection:Donor Org2	Char	7							Y
donorPathDesc3		Infection:Donor Org3	Char	7							Y
unitTestPerform	Y/N	Infection:Test to detect specific pathogen on unit post-Tx	Char	1							Y
unitTestPositive	Y/N	Infection:Unit - Positive/Negative results	Char	1							Y
unitPathDesc1		Infection:Unit Org1	Char	7							Y
unitPathDesc2		Infection:Unit Org2	Char	7							Y
unitPathDesc3		Infection:Unit Org3	Char	7							Y
nonirradBlood	Y/N	TA-GVND: Pt receive non-irradiated cellular product x 2mon preceding	Char	1							Y
unitHLA		TRALI: Donor or unit HLA specificity	Char	9							Y
	NOTEST	Not Done									





FieldName	Valid Values	Description	Data Type	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
deathdate		Death of death	Date	10	mm/dd/yyyy						Y
	01-12	Date of death -Month									
	01-31	Date of death-Day									
	2012-2020	Date of Death-Year									
x fusRelation		Relationship to death	Char	3							Y
	DEF	Definite									
	PRO	Probable									
	POS	Possible									
	DOU	Doubtful									
	RO	Ruled out									
	ND	Not determined									
unitImplicated		Unit Implicated in adverse reaction	Char	2		R					Y
	Y	Yes									
	N	No									
	NA	N/A									
numUnits		Total number of units transfused	Int	5		R					Y
compCode		Product Code	Char	5	Webform will allow 8-char Product Code.	R					Y
hemoCode		Check system used	Char	7		R					Y
	ISBT128	ISBT-128									
	CODABAR	Codabar									
unitNumber		Donation Identification Number (DIN) / Blood Unit Number	Char	16		R					Y
generalComment		General Comment	Char	3000							Y

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
EncounterID		EncounterID from Recipient database	Int	10							Y
SubjectID		SubjectID from Recipient database	Int	13							Y
rcpntPathDesc1OtherSpecified		Other organism entered (Y/N)	Char	1							Y
rcpntPathDesc2OtherSpecified		Other organism entered (Y/N)	Char	1							Y
rcpntPathDesc3OtherSpecified		Other organism entered (Y/N)	Char	1							Y
donorPathDesc1OtherSpecified		Other organism entered (Y/N)	Char	1							Y
donorPathDesc2OtherSpecified		Other organism entered (Y/N)	Char	1							Y
donorPathDesc3OtherSpecified		Other organism entered (Y/N)	Char	1							Y
unitPathDesc1OtherSpecified		Other organism entered (Y/N)	Char	1							Y
unitPathDesc2OtherSpecified		Other organism entered (Y/N)	Char	1							Y
unitPathDesc3OtherSpecified		Other organism entered (Y/N)	Char	1							Y

Specifications for RC\_VentilatorDuration

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
EncounterID		ID for Encounter. Will appear on all records in all tables for a specific encounter.	Int	10		R			Y	Y	Y
SubjectID		SubjectID will uniquely identify subject and will appear on all records in all tables.	Int	13		R			Y	Y	Y
StartDateD		Date ventilator started - Day	Int	2	dd	R			Y	Y	Y
	1-31										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
StartTime		Date ventilator started - Month	Int	2	mm	R			Y	Y	Y
	1-12										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
StopDateY		Date ventilator started - Year	Int	4	yyyy	R			Y	Y	Y
	2012-2020										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
StopDateD		Time ventilator started	Time	5	hh:mm	R			Y	Y	Y
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
		Date ventilator stopped - Day							Y	Y	Y
StopTime	1-31		Int	2	dd	R					
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									

FieldName	Valid Values	Description	DataType	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
StopDateM		Date ventilator stopped - Month	Int	2	mm	R			Y	Y	Y
	1-12										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
StopDateY		Date ventilator stopped - Year	Int	4	yyyy	R			Y	Y	Y
	2012-2020										
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
StopTime		Time ventilator stopped	Time	5	hh:mm	R			Y	Y	Y
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									
	-9	NOT COLLECTED at Hospital or Hub									
Duration		Duration of ventilator support (days). Calculated from date/time ventilator started and stopped.	Single	4.1	days				Y	Y	Y

Specifications for RC\_Vitals

FieldName	Valid Values	Description	Data Type	Length	Units	Required	LOD	Patient Subgroup (see text)			
								NR	TS	TCM	TX
EncounterID		ID for Encounter. Will appear on all records in all tables for a specific encounter.	Int	10		R		Y	Y	Y	Y
SubjectID		SubjectID will uniquely identify subject and will appear on all records in all tables.	Int	13		R		Y	Y	Y	Y
VitalSignType		Defines type of Vital sign.	Int	2		R		Y	Y	Y	Y
1	Height			Y	Y		Y	Y			
2	Weight			Y	Y		Y	Y			
3	Blood Pressure - Systolic			Y	Y		Y	Y			
4	Blood Pressure - Diastolic			Y	Y		Y	Y			
5	Mean Arterial Pressure			Y	Y		Y	Y			
6	Pulse			Y	Y		Y	Y			
7	Respiration			Y	Y		Y	Y			
8	Body Temperature			Y	Y		Y	Y			
9	CVP (Central Venous Pressure )/Right Atrial Pressure			Y	Y		Y	Y			
10	GCS (Glasgow Coma Scale for neurologic score)			Y	Y		Y	Y			
11	PCWP (Pulmonary Capillary Wedge Pressure)			Y	Y		Y	Y			
12	SpO2 (Pulse Oxygenation -O2 Saturation)			Y	Y		Y	Y			
Value		Vital Sign Value	Single	5.2		R		Y	Y	Y	Y
	-6	Other Notation									
	-8	NOT REPORTED or UNKNOWN for Participant									



Medication Codes for RC\_Medications table

Medication Code	Medication Name	Notes
10	3 Factor PCCs (prothrombin complex concentrates)	1/15/13, 7/31/13, 7/25/14
20	4 Factor PCCs (prothrombin complex concentrates)	
22	Albumin concentrate	3/12/2013: Added
30	Amicar (Epsilon amino caproic acid)	
40	Amiloride (Midamor)	
50	Apixaban	
60	Aspirin	
70	Bumetanide (Bumex)	corrected spelling 7/25/2014
80	Betrixaban	
90	Chlorothiazide (Diuril)	
100	Chlorthalidone (Hygroton)	
110	Coumadin	
120	Dabigatran	Corrected spelling 5/12/2015
130	Dalteparin	
140	Edoxaban	
150	Enoxaparin	
160	Eplerenone (Inspra)	
170	EPO (Erythropoietin)	
171	Darbepoetin	Addd 1/13/2015
180	Factor VIII	
190	Factor IX	
200	FEIBA	
210	Fondaparinux	
220	Furosemide (Lasix)	
230	Heparin	
240	Hydrochlorothiazide (Esidrix, Hydrodiuril)	

Medication Code	Medication Name	Notes
250	Humate-P	
260	Indapamide (Iozol)	
262	Intravenous immunoglobulin (IVIG)	Added 3/12/2013:
270	Metolazone (Zaroxolyn)	
280	Plavix	
282	Rh Immune Globin IM	Added 3/12/2013, Deleted 1/13/2015
283	Rh Immune Globin IV	Added 3/12/2013, Deleted 1/13/2015
284	Rh Immune Globin	Added 3/12/2013, Deleted 1/13/2015
290	Rivaroxaban	
300	Spiromolactone (Aldactone)	
310	Torsemide (Demadex)	
320	Triamterene (Dyrenium)	
330	TXA (tranexamic acid)	12/21/2015: corrected spelling
340	Vitamin K	
350	VIIa (Factor VIIa)	
99999	Not a primary med of interest	

Organisms for RC\_MicrobialTests

Organism Name	Organism Code	Notes
Acinetobacter species	10	
Aspergillus species	20	
Actinomyces species	30	
Aerococcus species	40	
Burkholderia species	50	
Candida species	60	
Citrobacter species	70	
Clostridia species	80	
Coagulase negative staphylococcus	90	
Corynebacterium species	100	
Eikenella species	110	
Enterobacter species	120	
Enterococcus species	130	
Escherichia coli	140	
Gram negative cocci	150	
Gram negative rods	160	
Gram positive cocci	170	
Gram positive rods	180	
Haemophilus species	190	
Klebsiella species	200	
Mucor species	210	
Mycobacterium species	220	
Neisseria species	230	
Normal Flora	240	
Peptostreptococcus species	250	
Propriionibacteria species	260	changed spelling from Propriionibacteria species to Propionibacteria species 4/1/2016
Propriionibacteria species	260	

<b>Organism Name</b>	<b>Organism Code</b>	<b>Notes</b>
Proteus species	270	
Pseudomonas species	280	
Serratia species	290	
Staphylococcus aureus	300	
Staphylococcus epidermidis	310	
Stenotrophomonas maltophilia	320	
Streptococcus pneumoniae	330	changed spelling from Streptococcus pneumoniae to Streptococcus pneumoniae 7/25/14
Streptococcus pneumoniae	330	
Streptococcus species	340	
Streptococcus viridans	350	
Other	360	

### Organisms for RC\_TxReactions

<b>Organism Name</b>	<b>Code</b>
<i>Actinomyces propionicus</i>	PRPRO
<i>Aerococcus christensenii</i>	AECH
<i>Aerococcus Genus</i>	AEGU
<i>Aerococcus sanguicola</i>	AESG
<i>Aerococcus sanguinicola</i>	AESGN
<i>Aerococcus spp.</i>	AESP
<i>Aerococcus urinae</i>	AEUR
<i>Aerococcus urinaeequi</i>	AEURQ
<i>Aerococcus urinaehominis</i>	AEURH
<i>Aerococcus viridans</i>	AEVI
<i>Arachnia propionica</i>	PRPRO
<i>Arthrobacter variabilis</i>	CORVAR
<i>Bacillus aeolius</i>	BAEOL
<i>Bacillus aerius</i>	BAERI
<i>Bacillus agaradhaerens</i>	BAGAR
<i>Bacillus alcalophilus</i>	BALCA
<i>Bacillus algicola</i>	BALGI
<i>Bacillus amyloliquefaciens</i>	BAMYL
<i>Bacillus aquimaris</i>	BAQUI
<i>Bacillus arseniciselenatis</i>	BARSE

<b>Organism Name</b>	<b>Code</b>
<i>Bacillus asahii</i>	BASAH
<i>Bacillus atrophaeus</i>	BATRO
<i>Bacillus azotoformans</i>	BAZOT
<i>Bacillus badius</i>	BBADI
<i>Bacillus barbaricus</i>	BBARB
<i>Bacillus bataviensis</i>	BBATA
<i>Bacillus benzoevorans</i>	BBENZ
<i>Bacillus boroniphilus</i>	BBORO
<i>Bacillus butanolivorans</i>	BBUTA
<i>Bacillus carboniphilus</i>	BCARB
<i>Bacillus cereus</i>	BC
<i>Bacillus cereus group</i>	BCERG
<i>Bacillus chittonlyticus</i>	BCHIT
<i>Bacillus circulans</i>	BACCIR
<i>Bacillus circulans group</i>	BACCIRG
<i>Bacillus clarkii</i>	BCLARK
<i>Bacillus clausii</i>	BCLAUS
<i>Bacillus coagulans</i>	BACCOA
<i>Bacillus cohnii</i>	BCOHN
<i>Bacillus decisifrondis</i>	BDECI
<i>Bacillus decolorationis</i>	BDECO

<b>Organism Name</b>	<b>Code</b>
Bacillus dipsosauri	BDIPS
Bacillus drentensis	BDREN
Bacillus edaphicus	BEDAP
Bacillus ehimensis	BEHIM
Bacillus endophyticus	BENDO
Bacillus farraginis	BFARR
Bacillus fastidiosus	BFAST
Bacillus firmus	BACFIR
Bacillus flexus	BFLEX
Bacillus fordii	BFORD
Bacillus fortis	BFORT
Bacillus fumarioli	BFUMA
Bacillus funiculus	BFUNI
Bacillus galactosidilyticus	BGALA
Bacillus gelatini	BGELA
Bacillus genus	BAC
Bacillus gibsonii	BGIBS
Bacillus ginsengi	BGINS
Bacillus globisporus ss. Marinus	BMARN
Bacillus gornadae	BGORN
Bacillus halmapalus	BHALM

<b>Organism Name</b>	<b>Code</b>
<i>Bacillus haloalkaliphilus</i>	BHALA
<i>Bacillus halodenitrificans</i>	BHALDF
<i>Bacillus halodurans</i>	BHALDR
<i>Bacillus halophilus</i>	BHALP
<i>Bacillus horikoshii</i>	BHORI
<i>Bacillus horti</i>	BHORT
<i>Bacillus humi</i>	BHUMI
<i>Bacillus hwajinpoensis</i>	BHWAJ
<i>Bacillus idriensis</i>	BIDRI
<i>Bacillus indicus</i>	BINDI
<i>Bacillus infantis</i>	BINFA
<i>Bacillus infernus</i>	BINFE
<i>Bacillus insolitus</i>	BINSO
<i>Bacillus jeotgali</i>	BJEOT
<i>Bacillus kaustophilus</i>	BKAUS
<i>Bacillus korlensis</i>	BKORL
<i>Bacillus krulwichiae</i>	BKRUL
<i>Bacillus laevolacticus</i>	BLAEV
<i>Bacillus lentinus</i>	BACLEN
<i>Bacillus licheniformis</i>	BACLIC
<i>Bacillus luciferensis</i>	BLUCI

<b>Organism Name</b>	<b>Code</b>
Bacillus macroides	BMACR
Bacillus macyae	BMACY
Bacillus marinus	BMARN
Bacillus marisflavi	BMARS
Bacillus massiliensis	BMASS
Bacillus megaterium	BACMEG
Bacillus methanolicus	BMETH
Bacillus mojavensis	BMOJA
Bacillus mucilaginosus	BMUCI
Bacillus muralis	BMURA
Bacillus mycoides	BMYCO
Bacillus naganoensis	BNAGA
Bacillus nealsonii	BNEAL
Bacillus niacini	BNIAC
Bacillus novalis	BNOVA
Bacillus odysseyi	BODYS
Bacillus okuhidensis	BOKUH
Bacillus oleronius	BOLER
Bacillus pallidus	BPALL
Bacillus pasteurii	BPAST
Bacillus patagoniensis	BPATA

<b>Organism Name</b>	<b>Code</b>
Bacillus pocheonensis	BPOCH
Bacillus pseudocaliphilus	BPSDL
Bacillus pseudofirmus	BPSDF
Bacillus pseudomycooides	BPSMY
Bacillus psychrodurans	BPSYD
Bacillus psychrophilus	BPSYP
Bacillus psychrosaccharolyticus	BPSYS
Bacillus psychrotolerans	BPSYT
Bacillus pulvifaciens	BPULV
Bacillus pumilus	BACPUM
Bacillus pycnus	BPYCN
Bacillus salexigens	BSALE
Bacillus saliphilus	BSALI
Bacillus schlegelii	BSCHL
Bacillus selenitireducens	BSELE
Bacillus shackletonii	BSHAC
Bacillus silvestris	BSILV
Bacillus simplex	BSIMP
Bacillus siralis	BSIRA
Bacillus smithii	BACSM
Bacillus soli	BSOLI

<b>Organism Name</b>	<b>Code</b>
Bacillus sonorensis	BSONO
Bacillus species not Bacillus anthracis	BACNANT
Bacillus sphaericus	BACSPH
Bacillus sporothermodurans	BSPOR
Bacillus spp.	BSP
Bacillus stearothermophilus	BACSTE
Bacillus subterraneus	BSUBT
Bacillus subtilis	BSU
Bacillus subtilis group	BACSUG
Bacillus subtilis spizizenii	BSSP
Bacillus subtilis ss. Inaquosorum	BSIN
Bacillus subtilis subtilis	BSS
Bacillus tequilensis	BTEQU
Bacillus thermantarcticus	BTHMN
Bacillus thermoamylovorans	BTHMM
Bacillus thermocatenulatus	BTHMC
Bacillus thermocloaceae	BTHMCL
Bacillus thermoglucoSIDASius	BTHMG
Bacillus thuringiensis	BACTHU
Bacillus tuscae	BTUSC
Bacillus vallismortis	BVALL

<b>Organism Name</b>	<b>Code</b>
<i>Bacillus vedderi</i>	BVEDD
<i>Bacillus velezensis</i>	BVELE
<i>Bacillus vietnamensis</i>	BVIET
<i>Bacillus vireti</i>	BVIRE
<i>Bacillus weihenstephanensis</i>	BWEIH
<i>Bacterionema matruchotii</i>	CORMA
<i>Brevibacterium ammoniagenes</i>	CORAMM
<i>Brevibacterium vitarumen</i>	CORVIT
<i>Corynebacterium accolens</i>	CORACC
<i>Corynebacterium acnes</i>	PRPAC
<i>Corynebacterium afermentans</i>	CORA
<i>Corynebacterium afermentans ss. Afermentans</i>	CORAA
<i>Corynebacterium afermentans ss. Lipophilum</i>	CORAL
<i>Corynebacterium ammoniagenes</i>	CORAMM
<i>Corynebacterium amycolatum</i>	CORAMY
<i>Corynebacterium anaerobium</i>	PRPAC
<i>Corynebacterium appendicis</i>	CORAPP
<i>Corynebacterium aquaticum</i>	CORAQ
<i>Corynebacterium aquilae</i>	CORAQL
<i>Corynebacterium argentoratense</i>	CORARG
<i>Corynebacterium atypicum</i>	CORATY

<b>Organism Name</b>	<b>Code</b>
<i>Corynebacterium aurimucosum</i>	CORAR
<i>Corynebacterium auris</i>	CORARS
<i>Corynebacterium auriscanis</i>	CORARSC
<i>Corynebacterium beticola</i>	CORBET
<i>Corynebacterium bovis</i>	CORBO
<i>Corynebacterium callunae</i>	CORCAL
<i>Corynebacterium camporealensis</i>	CORCAM
<i>Corynebacterium capitovis</i>	CORCAP
<i>Corynebacterium casei</i>	CORCAS
<i>Corynebacterium caspium</i>	CORCSP
<i>Corynebacterium ciconiae</i>	CORCIC
<i>Corynebacterium confusum</i>	CORCON
<i>Corynebacterium coyleae</i>	CORCOY
<i>Corynebacterium cystidis</i>	CORCY
<i>Corynebacterium diphtheroides</i>	PRPAC
<i>Corynebacterium durum</i>	CORDUR
<i>Corynebacterium efficiens</i>	COREFF
<i>Corynebacterium falsenii</i>	CORFAL
<i>Corynebacterium felinum</i>	CORFEL
<i>Corynebacterium flavescens</i>	CORFVS
<i>Corynebacterium flavidum</i>	CORST

<b>Organism Name</b>	<b>Code</b>
<i>Corynebacterium freneyi</i>	CORFRE
<i>Corynebacterium genitalium</i>	CORGEN
<i>Corynebacterium Genus</i>	CORGN
<i>Corynebacterium glaucum</i>	CORGLA
<i>Corynebacterium glucuronolyticum</i>	CORGLU
<i>Corynebacterium glutamicum</i>	CORGL
<i>Corynebacterium group A-3, CDC</i>	CORA3
<i>Corynebacterium group A-4, CDC</i>	CORA4
<i>Corynebacterium group A-5, CDC</i>	CORA5
<i>Corynebacterium group ANF, CDC</i>	CORA
<i>Corynebacterium group B-1, CDC</i>	CORB1
<i>Corynebacterium group B-3, CDC</i>	CORB3
<i>Corynebacterium group C, CDC</i>	CORC
<i>Corynebacterium group D-1, CDC</i>	CORD1
<i>Corynebacterium group D-2, CDC</i>	CORUR
<i>Corynebacterium group E, CDC</i>	CORE
<i>Corynebacterium group F-1, CDC</i>	CORF1
<i>Corynebacterium group F-2, CDC</i>	CORF2
<i>Corynebacterium group G-1, CDC</i>	CORACC
<i>Corynebacterium group G-2</i>	CORG2
<i>Corynebacterium group I-1, CDC</i>	CORI1

<b>Organism Name</b>	<b>Code</b>
<i>Corynebacterium</i> group I-2, CDC	CORI2
<i>Corynebacterium</i> group, CDC	CORG C
<i>Corynebacterium halotolerans</i>	CORHAL
<i>Corynebacterium hoagii</i>	CORHOA
<i>Corynebacterium hofmannii</i>	CORPD
<i>Corynebacterium imitans</i>	CORIMI
<i>Corynebacterium jeikeium</i>	CORJK
<i>Corynebacterium kroppenstedtii</i>	CORKRO
<i>Corynebacterium kutscheri</i>	CORKU
<i>Corynebacterium lilium</i>	CORGL
<i>Corynebacterium lipophiloflavum</i>	CORLIP
<i>Corynebacterium liquifaciens</i>	PRPAC
<i>Corynebacterium macginleyi</i>	CORMAC
<i>Corynebacterium manihot</i>	CORMAN
<i>Corynebacterium massiliense</i>	CORMSL
<i>Corynebacterium mastitidis</i>	CORMST
<i>Corynebacterium matruchotii</i>	CORMA
<i>Corynebacterium mediolanum</i>	CORMED
<i>Corynebacterium minutissimum</i>	CORMI
<i>Corynebacterium mooreparkense</i>	CORVAR
<i>Corynebacterium mucifaciens</i>	CORMUC

<b>Organism Name</b>	<b>Code</b>
<i>Corynebacterium murisepticum</i>	CORMRS
<i>Corynebacterium murium</i>	CORKU
<i>Corynebacterium mycetoides</i>	CORMY
<i>Corynebacterium nephridii</i>	CORNEP
<i>Corynebacterium nigricans</i>	CORAR
<i>Corynebacterium ovis</i>	CORPS
<i>Corynebacterium parvum</i>	PRPAC
<i>Corynebacterium phocae</i>	CORPHO
<i>Corynebacterium pilosum</i>	CORPI
<i>Corynebacterium propinquum</i>	CORPRO
<i>Corynebacterium pseudodiphtheriticum</i>	CORPD
<i>Corynebacterium pseudogenitalium</i>	CORPST
<i>Corynebacterium pseudotuberculosis</i>	CORPS
<i>Corynebacterium pseudotuberculostearicum</i>	CORPSM
<i>Corynebacterium renale</i>	CORRE
<i>Corynebacterium renale type 1</i>	CORRE
<i>Corynebacterium renale type II</i>	CORPI
<i>Corynebacterium renale type III</i>	CORCY
<i>Corynebacterium resistens</i>	CORRES
<i>Corynebacterium riegelii</i>	CORRIE
<i>Corynebacterium rubrum</i>	CORRUB

<b>Organism Name</b>	<b>Code</b>
<i>Corynebacterium seminale</i>	CORGLU
<i>Corynebacterium simulans</i>	CORSIM
<i>Corynebacterium singulare</i>	CORSIN
<i>Corynebacterium sphenisci</i>	CORSPI
<i>Corynebacterium spheniscorum</i>	CORSPM
<i>Corynebacterium spp.</i>	COS
<i>Corynebacterium spp. not Corynebacterium diphtheriae</i>	CORNCD
<i>Corynebacterium spp. not Corynebacterium jekeium</i>	CORNcj
<i>Corynebacterium striatum</i>	CORST
<i>Corynebacterium suicordis</i>	CORSUI
<i>Corynebacterium sundsvallense</i>	CORSUN
<i>Corynebacterium tenuis</i>	CORTEN
<i>Corynebacterium terpenotabidum</i>	CORTER
<i>Corynebacterium testudinoris</i>	CORTES
<i>Corynebacterium thomssenii</i>	CORTHO
<i>Corynebacterium tuberculostearicum</i>	CORTUB
<i>Corynebacterium ulcerans</i>	CORUL
<i>Corynebacterium urealyticum</i>	CORUR
<i>Corynebacterium ureicelerivorans</i>	CORURE
<i>Corynebacterium variabile</i>	CORVAR
<i>Corynebacterium viscosum</i>	CORVIS

<b>Organism Name</b>	<b>Code</b>
<i>Corynebacterium vitaeruminis</i>	CORVIT
<i>Corynebacterium xerosis</i>	CORXE
<i>Corynebacterium</i> , toxigenic	CORTOX
Diphtheroids	DIPTH
Diphtheroids, aerobic	DIPTHAE
Diphtheroids, anaerobic	DIPTHAN
<i>Micrococcus antarcticus</i>	MICANT
<i>Micrococcus epidermidis</i>	SE
<i>Micrococcus flavus</i>	MICLUT
<i>Micrococcus Genus</i>	MICG
<i>Micrococcus glutamicus</i>	CORGL
<i>Micrococcus luteus</i>	MICLUT
<i>Micrococcus lylae</i>	MICLYL
<i>Micrococcus lysodeikticus</i>	MICLUT
<i>Micrococcus nishinomiyaensis</i>	MICNIS
<i>Micrococcus pyogenes</i> var. <i>albus</i>	SE
<i>Micrococcus</i> spp.	MS
<i>Micrococcus</i> subgroup 3	STASA
<i>Pediococcus urinaeaequi</i>	AEURQ
<i>Peptococcus saccharolyticus</i>	STASL
<i>Propionibacterium acidipropionici</i>	PRPA

<b>Organism Name</b>	<b>Code</b>
<i>Propionibacterium acnes</i>	PRPAC
<i>Propionibacterium arabinosum</i>	PRPA
<i>Propionibacterium australiense</i>	PRPAU
<i>Propionibacterium avidum</i>	PROAV
<i>Propionibacterium cyclohexanicum</i>	PRPCY
<i>Propionibacterium freudenreichii</i>	PRPF
<i>Propionibacterium freudenreichii</i> ss. <i>Fredenreichii</i>	PRPFF
<i>Propionibacterium freudenreichii</i> ss. <i>Shermanii</i>	PRPFS
<i>Propionibacterium Genus</i>	PRPG
<i>Propionibacterium granulosum</i>	PROGR
<i>Propionibacterium granulosus</i>	CORC
<i>Propionibacterium innocuum</i>	PFI
<i>Propionibacterium jensenii</i>	PRPJE
<i>Propionibacterium lymphophilum</i>	PRPLY
<i>Propionibacterium microaerophilum</i>	PRPMI
<i>Propionibacterium pentosaceum</i>	PRPA
<i>Propionibacterium petersonii</i>	PRPJE
<i>Propionibacterium prionicus</i>	PRPPR
<i>Propionibacterium propionicum</i>	PRPRO
<i>Propionibacterium raffinosaceum</i>	PRPJE
<i>Propionibacterium rubrum</i>	PRPTH

<b>Organism Name</b>	<b>Code</b>
<i>Propionibacterium</i> spp.	PRSU
<i>Propionibacterium technicum</i>	PRPJE
<i>Propionibacterium thoenii</i>	PRPTH
<i>Propionibacterium zeae</i>	PRPJE
<i>Propioniferax innocus</i>	PFI
<i>Sarcina luteus</i>	MICLUT
<i>Staphylococcus albus</i>	SE
<i>Staphylococcus arlettae</i>	STAARL
<i>Staphylococcus auricularis</i>	STAAR
<i>Staphylococcus capitis</i>	STACS
<i>Staphylococcus capitis</i> ss. <i>Capitis</i>	STACC
<i>Staphylococcus capitis</i> ss. <i>Urealyticus</i>	STACU
<i>Staphylococcus caprae</i>	STACAP
<i>Staphylococcus carnosus</i>	STACAR
<i>Staphylococcus carnosus</i> ss. <i>Carnosus</i>	STACRC
<i>Staphylococcus carnosus</i> ss. <i>Utilis</i>	STACUT
<i>Staphylococcus chromogenes</i>	STACHR
<i>Staphylococcus coagulase negative</i>	CNS
<i>Staphylococcus cohnii</i>	STACO
<i>Staphylococcus cohnii</i> ss. <i>Cohnii</i>	STASCO
<i>Staphylococcus cohnii</i> ss. <i>Urealyticum</i>	STASUR

<b>Organism Name</b>	<b>Code</b>
<i>Staphylococcus condimenti</i>	STACON
<i>Staphylococcus epidermidis</i>	SE
<i>Staphylococcus epidermidis albus</i>	SE
<i>Staphylococcus epidermidis, elastase-producing strain</i>	STAEEES
<i>Staphylococcus equorum</i>	STAEQ
<i>Staphylococcus equorum ss. Equorum</i>	STAEE
<i>Staphylococcus equorum ss. Linens</i>	STAEL
<i>Staphylococcus felis</i>	STAFL
<i>Staphylococcus fleurettii</i>	STAFL
<i>Staphylococcus gallinarum</i>	STAGA
<i>Staphylococcus hemolyticus</i>	STAHA
<i>Staphylococcus hominis</i>	STAHO
<i>Staphylococcus hominis ss. Hominis</i>	STAHOME
<i>Staphylococcus hominis ss. Novobiosepticus</i>	STANOV
<i>Staphylococcus hyicus ss. Chromogenes</i>	STACHR
<i>Staphylococcus kloosii</i>	STAKLO
<i>Staphylococcus lentus</i>	STALE
<i>Staphylococcus lugdunensis</i>	STALU
<i>Staphylococcus muscae</i>	STAMUS
<i>Staphylococcus nepalensis</i>	STANEP
<i>Staphylococcus pasteurii</i>	STAPAS

<b>Organism Name</b>	<b>Code</b>
<i>Staphylococcus piscifermentans</i>	STAPIS
<i>Staphylococcus pulvereri</i>	STAVIT
<i>Staphylococcus saccharolyticus</i>	STASL
<i>Staphylococcus saprophyticus</i>	STASA
<i>Staphylococcus saprophyticus ss. Bovis</i>	STASB
<i>Staphylococcus saprophyticus ss. Saprophyticus</i>	STASAP
<i>Staphylococcus schleiferi</i>	STASH
<i>Staphylococcus schleiferi ss. Schleiferi</i>	STASCH
<i>Staphylococcus sciuri</i>	STASC
<i>Staphylococcus sciuri ss. Carnaticus</i>	STASCN
<i>Staphylococcus sciuri ss. Lentus</i>	STALE
<i>Staphylococcus sciuri ss. Rodentium</i>	STASRO
<i>Staphylococcus scuiri ss. Scuiri</i>	STASCU
<i>Staphylococcus simulans</i>	STASI
<i>Staphylococcus simulans biovar staphylolyticus</i>	STASBS
<i>Staphylococcus succinus</i>	STASU
<i>Staphylococcus succinus ss. Casei</i>	STASUC
<i>Staphylococcus succinus ss. Succinus</i>	STASUS
<i>Staphylococcus vitulinus</i>	STAVIT
<i>Staphylococcus vitulus</i>	STAVT
<i>Staphylococcus warneri</i>	STAWA

<b>Organism Name</b>	<b>Code</b>
<i>Staphylococcus xylosus</i>	STAXY
<i>Streptococcus anginosus milleri group</i>	STRANG
<i>Streptococcus anginosus</i>	STRVN
<i>Streptococcus anginosus group</i>	STRANG
<i>Streptococcus anginosus-constellatus</i>	STRAC
<i>Streptococcus australis</i>	STRAUS
<i>Streptococcus constellatus</i>	STRVC
<i>Streptococcus constellatus ss. Constellatus</i>	STRCC
<i>Streptococcus constellatus ss. Pharyngis</i>	STRCP
<i>Streptococcus criceti</i>	STRCRC
<i>Streptococcus cristatus</i>	STRCR
<i>Streptococcus dolonei</i>	STRD
<i>Streptococcus downei</i>	STRD
<i>Streptococcus entericus</i>	STRENT
<i>Streptococcus ferus</i>	STRF
<i>Streptococcus gordonii</i>	STRVG
<i>Streptococcus group F</i>	GFS
<i>Streptococcus infantis</i>	STRINF
<i>Streptococcus intermedius</i>	SVI
<i>Streptococcus milleri group</i>	STRANG
<i>Streptococcus milleri group A</i>	STRBA

<b>Organism Name</b>	<b>Code</b>
Streptococcus milleri group C	STRBC
Streptococcus milleri group F	GFS
Streptococcus milleri group G	STRBG
Streptococcus mitis	STRVM
Streptococcus mitis group	STRMIT
Streptococcus mutans	STRVT
Streptococcus mutans group	STRMUG
Streptococcus mutans serotype a	STRCRC
Streptococcus mutans serotype b	STRRT
Streptococcus mutans serotype h	STRD
Streptococcus mutans ss. Ferus	STRF
Streptococcus mutans ss. Sobrinus	STRSO
Streptococcus oralis	STROR
Streptococcus parasanguis	STRPA
Streptococcus peroris	STRPER
Streptococcus ratti	STRRT
Streptococcus salivarius	STRVS
Streptococcus salivarius group	STRSAG
Streptococcus salivarius ss. Salivarius	STRSLV
Streptococcus salivarius ss. Thermophilus	STRSAL
Streptococcus sanguis	SVS

<b>Organism Name</b>	<b>Code</b>
Streptococcus sanguis group	STRSG
Streptococcus sanguis II	STRSAN
Streptococcus sanguis type II	STRSAN
Streptococcus sobrinus	STRSO
Streptococcus thermophilus	STRSAL
Streptococcus vestibularis	STRVE
Streptococcus viridans group	SVU
Streptococcus, small-colony-forming beta-hemolytic group A	STRBA
Streptococcus, small-colony-forming beta-hemolytic group C	STRBC
Streptococcus, small-colony-forming beta-hemolytic group G	STRBG
Streptococcus, tufted fibril group	STRCR
Streptococcus, tufted mitior	STRSAN

Antibody List for RC\_Diagnostics

Antibody Name	Description	Notes
Anti-A		
Anti-A,B		
Anti-A1		
Anti-AUB		
Anti-B		
Anti-Bga		
Anti-Bgb		
Anti-Bgc		
Anti-C		
Anti-c		
Anti-Ce		
Anti-CHA		
Anti-COa		
Anti-COb		
Anti-CSa		
Anti-Cw		
Anti-D		
Anti-Dia		
Anti-Dib		
Anti-DOa		
Anti-DOB		
Anti-E		
Anti-e		
Anti-f		
Anti-FYa		
Anti-FYb		
Anti-G		

<b>Antibody Name</b>	<b>Description</b>	<b>Notes</b>
Anti-Ge		
Anti-Gya		
Anti-H		
Anti-HRb		
Anti-HY		
Anti-I		
Anti-i		
Anti-Jka		
Anti-JKb		
Anti-JMH		
Anti-JOa		
Anti-JSa		
Anti-JSb		
Anti-K		
Anti-k		
Anti-KNa		
Anti-KPa		
Anti-KPb		
Anti-LEa		
Anti-LEb		
Anti-LUa		
Anti-LUb		
Anti-LWa		
Anti-LWb		
Anti-M		Added 5/12/2015
Anti-M (IgG)		
Anti-M (IgM)		
Anti-McC		
Anti-N		

Antibody Name	Description	Notes
Anti-P		
Anti-P1		
Anti-RGa		
Anti-S		
Anti-s		
Anti-Sc1		
Anti-Sc2		
Anti-SDa		
Anti-SLA		
Anti-U		
Anti-V		
Anti-VEL		
Anti-WRa		
Anti-Xga		
Anti-YKa		
Anti-YTa		
Anti-YTb		
Cold Ab		
Cold auto Ab		
Fy3		
Passive D		
WRM	Warm Autoantibodies	
Other	"Other" should be used for any true, specific antibody not specified in list, including "Other"	
Non-Specific	If "Non-specific" is reported, then code as "Non-specific"	
-6	Everything else, including free text, such as "Specimen Possibly Contaminated. Results May Be Adversely Affected."	

Medications for medications table

Organisms for microbialtests

Organisms table from the txreactions spreadsheet for the txreactions table.

Antibody list for diagnostics table