Wisconsin Immunization Registry

HL7 – 2.3.1 – Implementation Guide for Immunization Messaging

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Wisconsin Immunization Registry

HL7 – General Transfer Specification (2.3.1)

Introduction

The Wisconsin Immunization Registry (WIR) has made available an interactive user interface on the World Wide Web for authorized users to enter, query, and update client immunization records. The Web interface makes WIR information and functions available on desktops around the state. However, some immunization providers already store and process similar data in their own information systems and may wish to keep using those systems while also participating in the statewide central repository. Others having billing needs and do not want to enter data into two diverse systems. WIR is being enhanced to allow providers to use the HL7 Version 2.3.1 to submit client and immunization information to WIR.

The Health Level Seven (HL7) Standard

The ANSI HL7 standard is widely used for data exchange in the health care industry. The full standard is quite lengthy, covering a variety of situations in patient care and health care finance, and no single application is likely to use all of its content. The CDC has worked with HL7 developers to create a set of messages that permit exchange of immunization data. This document covers the subset of HL7 that will be used for client and immunization records exchanged between WIR and outside systems.

- The basic unit transmitted in an HL7 implementation is the **message**.
- Messages are made up of several **segments**, each of which is one line of text, beginning with a three-letter code identifying the segment type.
- Segments are in turn made up of several **fields** separated by a delimiter character, "|".

MSH|^-\&|VALLEY CLINIC^^|||WIR^^\|19991005032342||VXU^V04|682299|P^\2.3.1^^\||ER PID||79928^^\|A5SMIT0071^^\\||SMITH^MARY^T^\\|JOHNSON^\\\|19951212|F|||| RXA|0|999|19970903|19970903|^\90701^DTP^CPT|0.5

The details of how HL7 messages are put together for WIR purposes will be explained later in this document; the example above shows the essentials of what a message looks like. In this example, a message is being sent on behalf of Valley Clinic to WIR consisting of three segments. NOTE: Valley Clinic may or may not be the actual transmitter of the message. The transmitter of the message will be identified by WIR from log-in information and not from an HL7 message.

- The Message Header segment (MSH) identifies the owner of the information being sent as (VALLEY CLINIC) and receiver (WIR) and identifies the message as being of type VXU, Unsolicited Vaccination Record Update, one of the message types defined by HL7.
- The Patient Identification segment (**PID**) gives the client's name (MARY T SMITH), birth date (19951212, in YYYYMMDD format), and other identifying fields.
- The Pharmacy Administration segment (**RXA**) tells that a DTP vaccine, with CPT code 90701, was administered on September 3, 1997 (formatted as 19970903). Many fields are optional, and this example could have included more information. Some segments may be repeated within a single message. In this example, the message could have included a second RXA segment to record another immunization given.

HL7 does not specify how messages are transmitted. It is flexible enough to be used for both real-time interaction and large batches. The standard defines file header and file trailer segments that are used when a number of messages are gathered into a batch for transmission as a file. WIR will use batch files of messages to communicate with outside systems.

Scope of This Document

The General Transfer Specification (GTS) documented here supports automated exchange of data between the WIR repository and outside systems, making client and immunization records available in both places while avoiding the need to enter data twice. The remainder of this document specifies how files of HL7 messages are constructed for WIR purposes. It does not cover the methods used to transmit files between the WIR central repository and outside systems. It covers only a small subset of the very extensive HL7 standard. Files of messages constructed from the guidelines in this document will fall within the HL7 standard, but there is a wide variety of other possible HL7 messages that are outside the scope of this document.

Disclaimer:

WIR's Web Service and PHIN-MS transports are designed for "real-time" single messaging. Organizations should avoid sending a cannonade (barrage) of messages to WIR at a single given instance. If you have a large volume of messages that you need processed, WIR requests that you create a batch file and submit them via WIR batch process.

References

- See Version 2.3.1 (June 1999) of the Health Level 7 standard for a full description of all messages, segments, and fields. Information regarding HL7 is at www.hl7.org.
- The National Immunization Program within the Center for Disease Control (www.cdc.gov/nip) has published an Implementation Guide for Immunization Data with the purpose of keeping the use of HL7 for immunization data as uniform as possible.

HL7 Message Types Used in WIR Transmissions

WIR uses messages of three types, one for sending client data without immunizations, one for sending immunizations, and one for acknowledging messages received. The tables below show how a message of each type is constructed from several segments. Each segment is one line of text, ending with the carriage return character, so HL7 messages are entirely readable and printable, though they may appear somewhat cryptic due to the scarcity of white space. (The standard has provisions for inclusion of binary data, but WIR will not use these features.) Square brackets [] enclose optional segments, and curly braces {} enclose segments that may be repeated. Thus, a message of type ADT could be composed of just MSH and PID segments, or these could be followed by one, two, or any number of NK1 segments. The full HL7 standard allows additional segments within these message types, but they are unused by WIR. In order to remain compliant with HL7, their use is not an error, but the message recipient can ignore their content. The segments documented here are sufficient to support the principal WIR functions of storing data about clients and immunizations.

Note: When sending messages to WIR, if your message contains segments that are NOT defined herein, your messages will <u>NOT</u> be rejected by WIR. In the event that your message contains extraneous segments, WIR will ignore the segment (and all corresponding datum values).

ADT

Update Patient Information

MSH Message Header
PID Patient Identification

[{NK1}] Next of Kin / Associated Parties

[{*OBX}] Observation/Result

VXU

Unsolicited Vaccination Record Update

MSH Message Header
PID Patient Identification

[PD1] Patient Additional Demographic
[{NK1}] Next of Kin / Associated Parties
{RXA} Pharmacy / Treatment Administration

[RXR] Pharmacy / Treatment Route (Only one RXR per RXA segment)

[{OBX}] Observation/Result*

ACK

General Acknowledgment
MSH Message Header

MSA Message Acknowledgment

[ERR] Error

*The only OBX segment valid within an ADT message is one indicating a CONTRAINDICATION specified in OBX-03 Value Type field. (e.g., 60009-8^Contraindication^LN)

RECOMMENDATIONS:

WIR will NOT accept an ADT message (unsolicited demographic update) for a new client unless at least ONE immunization exists for the client in WIR. Therefore, it is best to include the demographic update information in a VXU message whenever possible, as this message type accommodates BOTH immunization information and demographic update information. Should the provider wish to submit a new client using the ADT message, it must follow the VXU message for the new client within the file.

When a VXU^V04 (Unsolicited Vaccination Record Update) message type is sent with no RXA segment, a check is done to verify if the client exists in WIR or not. If the client already exists in WIR, then the demographic update will occur (*if all other update business rules apply). If the client is new to WIR, then the client will be rejected per current business rules.

Message Segments: Field Specifications and Usage

HL7 Segment Structure

Each segment consists of several fields, separated by the field separator character, "|". The tables below that define how each segment is structured contain the following columns:

1. SEQ	The ordinal position of the field in the segment. Since WIR does not use all possible fields in
	the HL7 standard, these are not always consecutive. When datum values are provided for fields
	NOT defined in this guide, WIR will ignore and NOT retain the datum value.

2. LEN Maximum length of the field

3. DT HL7 data type of the field. See below for definition of HL7 data types.

4. R/M R – required by HL7 M – mandatory for WIR Blank – optional field.

5. **RP**/# Y means the field may be repeated any number of times, an integer gives the maximum number of repetitions, and a blank means no repetition is permitted. WIR supports repetition

only for OBX-05.

6. TBL# Number of the table giving valid values for the field.

7. ELEMENT NAME HL7 name for the field.

- **HL7 data types.** Each field has an HL7 data type. Appendix A of this document lists and defines the HL7 data types needed for WIR. The elemental data types Numeric (NM) and String (ST) consist of one value, while some data types, such as Extended Person Name (XPN) are composites.
- **Delimiter characters.** Field values of composite data types consist of several components separated by the **component separator**, "^". When components are further divided into sub-components, these are separated by the **sub-component separator**, "&". Some fields are defined to permit repetitions separated by the **repetition character**, "~". When these special characters need to be included within text data, their special interpretations are prevented by preceding them with the **escape character**, "\".

MSH|^~\&|
XXX|field1|component1^component2^subcomponent3.1&subcomponent3.2^component4|
YYY|repetition1~repetition2|
ZZZ|data includes escaped \\~ special characters|

In the example above, the Message Header segment, as its definition requires, uses the field separator "i" immediately after the "MSH" code identifying the segment, and this establishes what character serves as the field separator throughout the message. The next field, the four characters "\~\&", establishes, in order, the component separator character, the repetition character, the escape character, and the sub-component separator character that will apply throughout the message. The hypothetical "XXX" segment includes field1 with no internal structure, but the next field has several components separated by "\", and the third of these is made up of two sub-components separated by "\". The hypothetical "YYY" segment's first field permits repetition, in this example the two values "repetition1" and "repetition2". The hypothetical "ZZZ" segment's field has a text value that includes the characters "\", and these are escaped to prevent their normal structural interpretation.

In WIR usage, sub-components, repetition, and text values requiring the escape character will be rare. Components within fields are common, since names and addresses are represented this way. HL7 permits use of other delimiters besides the recommended ones, and the delimiters used in each message are given in the Message Header segment. However, WIR will always use the recommended delimiters when sending files and requires their use for files received.

Rules for Sending Systems

The following rules are used by sending systems to construct HL7 messages.

- Encode each segment in the order specified in the message format.
- Begin the segment with the 3-letter segment ID (for example RXA).
- Precede each field with the data field separator ("|").
- Use HL7 recommended encoding characters ("^~\&").
- Encode the data fields in the order given in the table defining segment structure.
- Encode the data field according to its HL7 data type format.
- Do not include any characters for fields not present in the segment. Since later fields in the segment are encoded by ordinal position, fields that are not present do not reduce the number of field separators in the segment. For example,

when the second and third fields are not present, the field separators maintain the ordinal position of the fourth field: |field1||field4

- Data fields that are present but explicitly null are represented by empty double quotes "".
- Trailing separators may optionally be omitted. For example, |field1|field2||||| is equivalent to |field1|field2, when field3 and subsequent fields are not present.
- End each segment with the segment terminator (<u>always</u> the carriage return character ASCII Hex 0D followed by Hex 0A (carriage-return and linefeed), or a single character Hex 0A (line feed).

Rules for Receiving Systems

The following rules are used by receiving systems to process HL7 messages.

- Treat data segments that are expected but not present as if all data fields in the segment were not present.
- Require use of HL7 recommended Field Separator |, and Encoding characters ^~\& for encoding messages.
- Ignore any data segment that is included but not expected, rather than treating it as an error. The HL7 message types used by WIR may include many segments besides the ones in this document, and WIR ignores them. WIR will not send messages with segments not documented in this specification, but reserves the right to specify more segments at a later date. The rule to ignore unexpected segments facilitates this kind of change.
- Ignore data fields found but not expected within a segment.

The message segments below are the ones needed to construct messages of the types used by WIR. Each segment is given a brief description excerpted from the HL7 standard. The tables define what fields make up each segment. Since WIR does not use all the fields HL7 defines, there are sometimes gaps in the ordinal sequence of fields. Following HL7 rules, the gaps do not diminish the number of field separators within the segment. For example, if the second and third fields in a segment are not present, their field separators remain in order to indicate that the next field present is the fourth: field1|||field4.

ERR

The ERR segment is used to add error comments to acknowledgment messages.

SEQ	LEN	DT	R/M	RP/#	TBL#	ELEMENT NAME
1	80	CM	R	Υ		Error Code and Location

Field Notes:

ERR-1 A composite field with four components.

<segment ID (ST)>^<sequence (NM)>^<field position (NM)>^<field sub-component ordinal number (NM)</pre>

The first component identifies the segment ID containing the error. The second component identifies the input file line number of the segment containing the error. The third component identifies by ordinal number the field containing the error. The fourth component identifies by ordinal number the field sub-component containing the error (0 if not applicable) The remaining five components of the CE data type are not valued and their '^' separators are not generated. Note that error text is transmitted in field MSA-3. For example, if the NK1 segment is missing a mandatory field:

ERR|NK1^10^2^1

This error message identifies the NK1 segment occurring on line 10 of the input file whose mandatory second field (Name) is missing the mandatory 1st component (Family Name).

MSA

The MSA segment contains information sent while acknowledging another message.

SEQ	LEN	DT	R/M	RP/#	TBL#	ELEMENT NAME
1	2	ID	R		8000	Acknowledgment Code
2	20	ST	R			Message Control ID
3	80	ST				Text Message

- MSA-1 Acknowledgement code giving receiver's response to a message. AA (Application Accept) means the message was processed normally. AE (Application Error) means an error prevented normal processing. An error message will be put in MSA-3, and for ACK messages the optional ERR segment will be included.
- MSA-2 The message control ID from MSH-10 in the message being acknowledged. This allows the sending system to associate this response with the message being responded to.
- MSA-3 Text of error message, used when MSA-1 does not have the normal value of AA.

MSH

The MSH segment defines the intent, source, destination, and some specifics of the syntax of a message.

SEQ	LEN	DT	R/M	RP/#	TBL#	ELEMENT NAME
1	1	ST	R			Field Separator
2	4	ST	R			Encoding Characters
3	180	EI				Sending Application
4	180	EI				Sending Facility
6	180	EI				Receiving Facility
7	26	TS				Date/Time Of Message
9	7	CM	R			Message Type
10	20	ST	R			Message Control ID
11	3	PT	R		0103	Processing ID
12	60	VID	R		0104	Version ID
15	2	ID			0155	Accept Acknowledgment Type

Field Notes:

- MSH-1 Determines the field separator in effect for the rest of this message. WIR requires the HL7 recommended field separator of "|".
- MSH-2 Determines the component separator, repetition separator, escape character, and sub-component separator in effect for the rest of this message. WIR requires the HL7 recommended values of ^~\&.
- MSH-3 Name of the sending application. When sending, WIR will use "WIR" followed by the current version number of the registry. This field is an optional convenience. See MSH-4 and MSH-6 for the fields principally used to identify sender and receiver of the message.
- MSH-4 Identifies for whom the message is being sent (the owner of the message information). When sending, WIR will use "WIR". When the message is being sent to WIR and the Provider Organization owning the information is different than the organization transmitting the message, use the WIR Provider ID of the Provider Organization that owns the information. Contact the WIR Help Desk for the appropriate organization ID. If the owner of the information and the transmitter of the information are the same Provider Organization, this field may be left blank.

This field is required for:

- 1. Providers that are sending via PHIN-MS or Web Services
- MSH-6 Identifies the message receiver. When sending, WIR will use the brief Provider Organization name assigned when the provider first registers with the WIR database and WIR-Web interface.
- MSH-7 Date and time the message was created. WIR ignores any time component. See the TS data type.
- MSH-9 This is a required field. Two components of this field give the HL7 message type (see Table 0076) and the HL7 triggering event (see Table 0003). Within HL7, the triggering event is considered to be the real-world circumstance causing the message to be sent. For WIR purposes, this field should have the value ADT^A31 for a message conveying client information or the value VXU^V04 for a message conveying client and immunization information. In acknowledgement messages the value ACK is sufficient and the second component may be omitted.
- MSH-10 This is a required field. The message control ID is a string (which may be a number) uniquely identifying the message among all those ever sent by the sending system. It is assigned by the sending system and echoed back in the ACK message sent in response.
- MSH-11 The processing ID to be used by WIR is **P** for production processing. If this field is null, an informational message is generated indicating that WIR is defaulting to **P**.
- MSH-12 This is a required field. Use a value of "2.3.1" to indicate HL7 Version 2.3.1.
- MSH-15 This field controls whether an acknowledgement is generated for the message sent. WIR suggests a value of ER to ask that acknowledgements be sent only for messages that cannot be processed normally. If the field is empty, WIR will assume the value of ER.

PID

The PID segment is used by all applications as the primary means of communicating patient identification information. This segment contains permanent patient identifying and demographic information that, for the most part, is not likely to change frequently.

SEQ	LEN	DT	R/M	RP/#	TBL#	ELEMENT NAME
2	20	CX				Patient ID (External ID)
3	20	CX	R			Patient ID (Internal ID)
5	48	XPN	R	Υ		Patient Name
6	48	XPN		Υ		Mother's Maiden Name
7	26	TS	М			Date/Time of Birth
8	1	IS			0001	Sex
10	80	CE		Υ	0005	Race
11	106	XAD				Patient Address
19	16	ST				SSN Number – Patient
22	80	CE		Υ	0189	Ethnic Group
24	1	ID			0136	Multiple Birth Indicator
25	2	NM				Birth Order
29	26	TS				Patient Death Date and Time

- PID-2 When a Provider Organization is sending to WIR, use the client's WIR Client ID if available. When WIR is sending to an outside system, it will use that system's Chart Number or other identifier if available. If a
- PID-3 When a Provider Organization is sending to WIR, use the sending system's Chart Number or other identifier if available. When WIR is sending to an outside system it will use the client's WIR ID and chart number when it is available. WIR does not support repetition of this field.
- PID-5 See the XPN data type. Last name and first name are required in the first two components. WIR does not support repetition of this field.
- PID-6 See the XPN data type. In this context, where the mother's name is used for client identification, WIR uses only last name and first name. A mother's legal name might also appear in the context of an NK1 segment. WIR does not support repetition of this field.
- PID-7 Give the year, month, and day of birth (YYYYMMDD). WIR ignores any time component.
- PID-8 See Table 0001. Use F, M, or U.
- PID-10 See Table 0005. WIR stores and writes "unknown" values as null. WIR does not accept Hispanic or Latino as a race option. Submit it in the Ethnic Group PID-22 field. WIR does not support repetition of this field.
- PID-11 See the XAD data type. WIR does not support repetition of this field.
- PID-19 NOTE: Social security number is used for identification purposes only, and is not displayed in screens or distributed to Provider Organizations.
- PID-22 See Table 0189. WIR does not support repetition of this field.
- PID-24 Use Y to indicate that the client was born in a multiple birth.
- PID-25 Relevant when client was born in a multiple birth. Use 1 for the first born, 2 for the second, etc. This field is useful in matching client data to existing records.
- PID-29 The date of death, if client is deceased. Give the year, month, and day (YYYYMMDD). WIR ignores any time component. If a death date is sent, then the Patient Registry Status in PD1-14 must indicate a value of "P" for permanently inactive/deceased.

PD1

The PD1 carries patient additional demographic information that is likely to change.

SEQ	LEN	DT	R/M	RP/#	TBL#	ELEMENT NAME
11	80	CE			0215	Publicity Code
12	1	ID			0136	Protection Indicator
14	40	IS			NIP006	Patient Registry Status

Field Notes:

- PD1-11 Controls whether recall/reminder notices are sent. WIR will recognize "01" to indicate no recall/reminder notices or "02" recall/reminder notices any method.
- PD1-12 Controls visibility of records to other organizations.
- PD1-14 See table NIP006.

NK1

The NK1 segment contains information about the patient's other related parties. Any associated parties may be identified. Utilizing *NK1-1-set ID*, multiple NK1 segments can be sent to patient accounts.

SEQ	LEN	DT	R/M	RP/#	TBL#	ELEMENT NAME
1	4	SI	R			Set ID – NK1
2	48	XPN		Υ		Name
3	60	CE			0063	Relationship
4	106	XAD		Υ		Address
5	40	XTN		Υ		Phone Number
22	80	CE			0215	Publicity Code

- NK1-1 Sequential numbers. Use "1" for the first NK1 within the message, "2" for the second, and so forth. Although this field is required by HL7, WIR will ignore its value, and there is no requirement that the record for the same responsible person keep the same sequence number across multiple messages, in the case that information from the same record is transmitted more than once.
- NK1-2 Name of the responsible person who cares for the client. See the XPN data type. WIR does not support repetition of this field.
- NK1-3 Relationship of the responsible person to the client. See data type CE and Table 0063 in the HL7 tables. Use the first three components of the CE data type, for example |32^Mother^HL70063^^^|.
- NK1-4 Responsible person's mailing address. See the XAD data type. WIR does not support repetition of this field.
- NK1-5 Responsible person's phone number. Format as (999)999-9999. WIR does not support repetition of this field.
- NK1-22 Controls whether recall/reminder notices are sent for the responsible person. WIR will recognize "01" to indicate no recall/reminder notices or "02" recall/reminder notices any method.

RXA

The RXA carries pharmacy administration data. It is a repeating segment and can record unlimited numbers of vaccinations.

SEQ	LEN	DT	R/M	RP/#	TBL#	ELEMENT NAME
1	4	NM	R			Give Sub-ID Counter
2	4	NM	R			Administration Sub-ID Counter
3	26	TS	R			Date/Time Start of Administration
4	26	TS	R			Date/Time End of Administration
5	100	CE	R			Administered Code
6	20	NM	R			Administered Amount
7	60	CE	С			Administered Units
9	200	CE		Υ	NIP001	Administration Notes
10	200	XCN				Administering Provider
11	200	CM				Administered-at location
15	20	ST				Substance Lot Number
17	60	CE			0227	Substance Manufacturer Name
18	200	CE			NIP002	Substance Refusal Reason
21	2	ID				Action Code-RXA

Field Notes:

- RXA-1 Required by HL7. Use "0" for WIR.
- RXA-2 Required by HL7. Use "999" for WIR
- RXA-3 Date the vaccine was given. WIR ignores any time component.
- RXA-4 Required by HL7. Ignored by WIR, which will use the value in RXA-3.
- RXA-5 This field identifies the vaccine administered. WIR accepts the CVX code, CPT code, Vaccine Trade Name, or Vaccine Group Code for the vaccine administered. If using the CVX code, give the CVX code in the first component and "CVX" in the third component. If using the CPT code, the vaccine group code or vaccine trade name, use components four through six. For example, give the CPT code in the fourth component and "CPT" in the sixth component, |^^90700^DtaP^CPT|. If using vaccine group code, use "WVGC" as the name of the coding system. If using vaccine trade name, use "WVTN" as the name of the coding system. See the CE data type and HL7 Table 0292 (CVX Codes), WIR Table WCPT (CPT Codes), WIR Table WVGC (Vaccine Group Codes), and WIR Table WVTN (Vaccine Trade Names).

RXA-6 When RXA-7 is not valued

This field value will be interpreted as Dose Magnitude – the number of age appropriate doses administered. For example, a dose magnitude of 2 of a pediatric formulation would be adequate for an adult. WIR and HL7 require this field to contain a value. However, a value of 1.0 will be stored in its place

When RXA-7 is valued

This field value will be interpreted as the dosage amount (e.g., 0.5, 0.65, 1.0, 1.5, etceteras.). The dose amount provided will be saved and displayed/reported.

- RXA-7 WIR will recognize any value to indicate that RXA-06 should be interpreted as the dosage amount. WIR will treat the immunization as 1 FULL dose and store and display/report administered unit (ML, gm, grams, CAP, etc.) that is provided.
- RXA-9 WIR will recognize 00 to indicate Administered Vaccine, 01 to indicate Historical Record or 07 to indicate School Record. When sending, WIR will include the corresponding immunization id in the second repeating segment.

|01^^^^~9999999WIR immunization id^IMM ID^^^|

The 07 value can only be used by organizations that are set up to send school information, otherwise the incoming immunization will be rejected.

- RXA-10 Identifies the name of the person physically administering the vaccine (the vaccinator). WIR will use components 2 7 to record the name and does not support repetition of this field.
- RXA-11 WIR will use this field to identify the facility where the vaccine was administered. Place the facility name in component 4.
- RXA-15 Manufacturer's lot number for the vaccine. WIR does not support repetition of this field.

- RXA-17 Vaccine manufacturer from Table 0227, for example |AB^Abbott^ MVX^^^|. The HL7 2.3.1 specification recommends use of the external code set MVX. "When using this code system to identify vaccines, the coding system component of the CE field should be valued as "MVX" not as "HL70227." WIR does not support repetition of this field
- RXA-18 When applicable, this field records the reason the patient refused the vaccine. See table NIP002. Any entry in this field indicates that the patient did not take the substance. The vaccine that was offered should be recorded in RXA-5, with the number 0 recorded for the dose number in RXA-2. Do not record contraindications, immunities or reactions in this field. WIR does not support repetition of this field.

Notes on Refusals:

- 6) WIR only stores the fact that a refusal of a vaccine occurred, not a specific type of refusal, so all outgoing refusals will be designated as "PARENTAL DECISION." Please see the example below.
- b) The WIR system will not write out refusals which do not have an applies-to date. It will write out multiple refusals for the same vaccine on different dates for those clients who have them.
- c) The WIR system will accept incoming refusals of the same vaccine on different dates and file them both. However, if they both have the same applies-to date, then only one will be stored.
- d) The sending organization will become the refusal owner. In general, only the organization who owns the refusal is permitted to edit it. However, in the case of parent and child organizations, the parent may edit the child's refusals and vice versa.

Here is a sample RXA segment for an MMR refusal given on the date 01/01/2007: RXA|0|0|20070101|20070101|^^^MMR^MVGC|1.0|||||||||00^PARENTAL REFUSAL^NIP002^^^

RXA-21 To delete an existing immunization in WIR specify a value of "D". In addition to requiring that the existing immunization is owned by the same provider requesting the delete, WIR limits that no more than 5% of all incoming immunizations can be flagged as delete and no more than 50 total.

RXR

The Pharmacy/Treatment Route Segment contains the alternative combination of route and site.

SEQ	LEN	DT	R/M	RP/#	TBL#	ELEMENT NAME
1	60	CE	R		0162	Route
2	60	CE			0163	Site

Field Notes:

- RXR-1 This is the route of administration from table 0162.
- RXR-2 This is the site of the route of administration from table 0163.

OBX

The Observation/Result Segment is used to transmit an observation.

SEQ	LEN	DT	R/M	RP/#	TBL#	ELEMENT NAME
1	10	SI				Set ID-OBX
2	3	ID				Value type
3	590	CE	R			Observation Identifier
5	65536	-	R	Υ		Observation Value
11	1	ID	R		0085	Observation Result Status
14	26	TS				Date/Time of the observation

- OBX-1 Sequential numbers. Use "1" for the first OBX within the message, "2" for the second, and so forth.
- OBX-2 This field contains the data type which defines the format of the observation value in OBX-5. For incoming PO-WIR data, Data Exchange accepts CE for Coded Entry. However, for WIR-PO, the system will send out values of CE, TS, NM for Coded Entry, Timestamp, and Number respectively, depending on what is actually sent in OBX-5.

For school data exchange, the system will also accept and send the value ID in OBX-2

OBX-3 When indicating a **Vaccination Contraindication/Precaution**, use 60009-8 in this field and enter a Contraindication, Precaution, or Immunity code (NIP004) in OBX-5.

When indicating a **Reaction to Immunization**, use 31044-1 in this field and enter a Reaction code (WIR001) in OBX-5.

Example: OBX|1|CE|31044-1^Reaction^LN||^HYPOTON^hypotonic^WIR^^^||||||F|

When indicating a **Vaccination Adverse Event Outcome**, use 60012-2 in this field and enter an Event Consequence code (NIP005) in OBX-5.

Example: OBX|1|CE|60012-2^Adverse Outcome^LN||^E^er room^NIP^^^||||||F|

When indicating a **FERPA Release Status**, use FERPA in this field and enter a Yes/No Indicator code (HL70136) in OBX-5.

Example: OBX|1|ID|FERPA^FERPA Release^99W01||Y||||||F|

When indicating a **Graduation Year**, use GRADYEAR in this field and enter a four digit year (YYYY) in OBX-5. Example: OBX|1|TS|GRADYEAR^Graduation Year^99W01||2023||||||F|

When indicating **Date Enrolled in WI School,** use ENROLLDATE in this field and give the year, month, and day that the student was first enrolled in Wisconsin Schools (YYYYMMDD) in OBX-5.

Example: OBX|1|TS|ENROLLDATE^Date Enrolled in WI School^99W01||20010825|||||||F|

OBX-5 Text reporting Contraindication, Precaution, or Immunity (NIP004), Reaction (WIR001), Event Consequence (NIP005), or WIR Student Information (99W01). WIR has imposed a CE data type upon this field. The first component of which is required.

(e.g., |PERTCONT^Pertussis contra^WIR^^^|)

- OBX-11 Required for HL7. Use "F" for WIR.
- OBX-14 Records the time of the observation. WIR ignores any time component.
- **NOTE 1:** The only valid OBX Observation Identifier (OBX-03) for an **ADT^A31** message type is Contraindication/Precaution (60009-8).
- **NOTE 2:** All OBX messages with an observation identifier of Vaccination Contraindication/Precaution will be returned in an outgoing file in a separate ADT message for the client.

Batch Files of HL7 Messages

The definitions above tell how to create messages containing client and immunization data. Each message can logically stand on its own, and HL7 is compatible with various methods of online and batch transmission. WIR uses batch files to transmit many messages together. HL7 provides special header and footer segments to structure batch files. These segments are not part of any message, but serve to bracket the messages defined above. The structure of a batch file is as follows.

FHS

File Header Segment

The FHS segment is used to head a file (group of batches).

SEQ	LEN	DT	R/M	RP/#	TBL#	ELEMENT NAME
1	1	ST	R			File Field Separator
2	4	ST	R			File Encoding Characters
3	15	ST				File Sending Application
4	20	ST	М			File Sending Facility
6	20	ST	М			File Receiving Facility
7	26	TS	М			File Creation Date/Time
9	20	ST	М			File Name/ID
10	80	ST				File Header Comment
11	20	ST	М			File Control ID
12	20	ST				Reference File Control ID

- FHS-1 Same definition as the corresponding field in the MSH segment.
- FHS-2 Same definition as the corresponding field in the MSH segment.
- FHS-3 Same definition as the corresponding field in the MSH segment.
- FHS-4 Same definition as the corresponding field in the MSH segment.
- FHS-6 Same definition as the corresponding field in the MSH segment.
- FHS-7 Same definition as the corresponding field in the MSH segment.
- FHS-9 Name of the file as transmitted from the initiating system.
- FHS-10 Free text, which may be included for convenience, but has no effect on processing.
- FHS-11 This field is used to identify a particular file uniquely among all files sent from the sending facility identified in FHS-4.
- FHS-12 Contains the value of FHS-11-file control ID when this file was originally transmitted. Not present if this file is being transmitted for the first time.

FTS

File Trailer Segment

The FTS segment defines the end of a file.

ĺ	SEQ	LEN	DT	R/M	RP/#	TBL#	ELEMENT NAME
	1	10	NM	М			File Batch Count
	2	80	ST				File Trailer Comment

Field Notes:

FTS-1 The number of batches contained in this file. WIR normally sends one batch per file and discourages sending multiple batches per file.

FTS-2 Free text, which may be included for convenience, but has no effect on processing.

BHS

Batch Header Segment

The BHS segment defines the start of a batch.

SEQ	LEN	DT	R/M	RP/#	TBL#	ELEMENT NAME
1	1	ST	R			Batch Field Separator
2	4	ST	R			Batch Encoding Characters
3	15	ST				Batch Sending Application
4	20	ST	М			Batch Sending Facility
6	20	ST	М			Batch Receiving Facility
7	26	TS	М			Batch Creation Date/Time
10	80	ST				Batch Comment
11	20	ST	М			Batch Control ID
12	20	ST				Reference Batch Control ID

Field Notes:

- BHS-1 This field contains the separator between the segment ID and the first real field, *BHS-2-batch encoding characters*. As such it serves as the separator and defines the character to be used as a separator for the rest of the segment. WIR requires | (ASCII 124).
- BHS-2 This field contains the four characters in the following order: the component separator, repetition separator, escape characters, and sub-component separator. WIR requires ^~\&, (ASCII 94, 126, 92, and 38, respectively).
- BHS-3 Same definition as the corresponding field in the MSH segment.
- BHS-4 Same definition as the corresponding field in the MSH segment.
- BHS-6 Same definition as the corresponding field in the MSH segment.
- BHS-7 Same definition as the corresponding field in the MSH segment.
- BHS-10 Free text, which may be included for convenience, but has no effect on processing.
- BHS-11 This field is used to uniquely identify a particular batch. It can be echoed back in *BHS-12-reference batch control ID* if an answering batch is needed. For WIR purposes, the answering batch will contain ACK messages.
- BHS-12 This field contains the value of *BHS-11-batch control ID* when this batch was originally transmitted. Not present if this batch is being sent for the first time. See definition for *BHS-11-batch control ID*.

BTS

Batch Trailer Segment

The BTS segment defines the end of a batch.

SEQ	LEN	DT	R/M	RP/#	TBL#	ELEMENT NAME
1	10	ST	М			Batch Message Count
2	80	ST				Batch Comment

Field Notes:

BTS-1 This field contains the count of the individual messages contained within the batch.

BTS-2 Free text, which may be included for convenience, but has no effect on processing.

File Interchange between WIR and Outside Systems

The central repository of WIR contains records of clients from around the state. Client and immunization records flow both ways between WIR and outside systems. Data for a particular client is transmitted by WIR to an outside system at a Provider Organization only if the client is already identified as having a relationship with that Provider Organization, and this relationship is created by transmitting the client's record to WIR. So an exchange through this General Transfer Specification of information about a given client is always initiated by the outside system. There are three (3) options for exchanging data with WIR. (1) The Provider Organization can send data to WIR and request no data be returned from WIR. (2) The Provider Organization can request data from WIR while not providing data to WIR. (3) The Provider Organization can send data to WIR and WIR will return any updated information regarding clients having a relationship with the Provider Organization to the Provider Organization.

Note that client and immunization data can also be entered, queried up, and modified using the WIR-Web interface, and this provides an alternate way of identifying a client as having a relationship with a Provider Organization. But use of WIR-Web is not required to create a relationship between a Provider Organization and a client, and the first transmission to WIR of a client immunization record creates the link that thereafter causes WIR to transmit that client's record to the outside system.

HL7 messages are always part of a two-way exchange between an initiating system and a responder. Sometimes the initial message implies specific data to be sent in a response. Other times, as is the case with WIR client and immunization data, the principal response of the receiving system is to process the message and post whatever it contains to its own database. For these cases, HL7 provides the ACK message type, which contains no new application data, but allows the receiver to inform the initiator that the message has been received and processed successfully. In case of an error that prevents successful processing, optional parts of the ACK message allow this to be communicated as well.

For exchanges between WIR and outside systems, it is the responsibility of the outside system to initiate the transfer of the first file, containing ADT and/or VXU messages with client and immunization data. After processing those messages, WIR responds with a file of ACK messages. At the same time or soon after, WIR also creates another file of ADT and VXU messages, containing the full client record, to send to the Provider Organization that initiated the first transfer. It is the responsibility of that Organization as receiver to transmit back a file of ACK messages. During this second exchange, in terms used by HL7, WIR is the initiator and the outside system is the respondent. However, it is the receipt of the first file initiated by the outside system that causes WIR to initiate sending its own data file.

	Provider Organization		WIR
		Outgoing	Receiving
1.	Creates a file of client and immunization records that have changed since they were last transmitted to WIR.		
2.	Transmits the file to WIR.		
3.			Processes the file received, creates a file of ACK messages.
4.		Transmits the ACK file back to the initiator of the original file.	
5.	Processes the ACK file to confirm success of the file transmission.		
6.			d immunization records that have last transmitted to this Provider
7.		Transmits this file to the Provider Organization.	
8.	Processes the file received, creates a file of ACK messages.		
9.	Transmits the ACK file back to WIR		
10.			Processes the ACK file to confirm success of the file transmission.

The 15th field in the MSH message header segment allows the initiator to ask that the message be acknowledged only in case of an error, and WIR suggests this choice to minimize the number of ACK messages transmitted. In this case the ACK file contains only error messages (an optional form of the ACK message type), and the original messages with no answering error messages are implicitly acknowledged as successfully processed. If all messages in a batch are successful, the answering ACK

file may contain only file and batch headers and footers, with no actual ACK messages. In Step 1 in the above table, it is permissible for a Provider Organization to send a file containing only file and batch headers and footers as a way of triggering the file WIR creates in Step 6. It is also possible for the file WIR creates in Step 6 to contain only file and batch headers and footers, if there are no records selected to send.

Examples

To illustrate how a WIR HL7 file is put together, we will show how the fictional Valley Clinic formats client and immunization records to transmit to WIR. The following table shows the information to be transmitted, organized into HL7 segments and fields. For example, PID-3 refers to the third field in the Patient Identification segment.

Information to transmit	Data value to be entered	HL7 Format
	(See page 13 for format)	
• Client #1		PID segment
Chart Number (ID on Valley Clinical autom)	45LR999	PID-3
Clinic's system)	GEORGE M MILLER JR	PID-5
Name Mather's maiden name	MARTHA OLSON	PID-5
Mother's maiden name Pinth data	February 27, 1995	PID-0
Birth date	M	PID-7
SexAddress	123 MAIN ST	PID-11
	MADISON, WI 53000, WI025	
 Social Security Number 	000111222	PID-19
Multiple Birth Indicator	Y (client was born as part of a multiple birth)	PID-24
Birth Order	2 (second birth of a multiple birth)	PID-25
Publicity Code	02	PD1-11
Protection Indicator	Y (client records are visible by other provider organizations)	PD1-12
Patient Registry Status	A (client is active in the registry)	PD1-14
Responsible Person (parent or other person who cares for client)		NK1 segment
Name	MARTHA MILLER	NK1-2
Relationship to client	32	NK1-3
• Address	123 MAIN ST MADISON, WI 53000, W1025	NK1-4
• Phone	608 123 4567	NK1-5
Responsible Person		NK1 segment
Name	GEORGE MILLER	NK1-2
Relationship to client	33	NK1-3
Responsible Person		NK1 segment
Name	LUCAS JONES	NK1-2
Relationship to client	D3	NK1-3
• Address	MADISON, WI 53715	NK1-4
• Phone	515 829 1521	NK1-5
Publicity Code	02	NK1-22
• Client #2		PID segment
WIR Client ID (Valley Clinic received this in an earlier transmission from WIR)	66782	PID-2
Chart Number	23LK729	PID-3
Name	MARIA CALIFANO	PID-5
Mother's maiden name	ANGELICA DISTEFANO	PID-6
Birth date	April 13, 1998	PID-7
• Sex	F	PID-8
• Immunization		RXA segment

Date administered	July 23, 1999	RXA-3
	DtaP	RXA-5
	90700	RXA-5
• CPT Code		
• Dose (ml)	0.5	RXA-6
 Administering Provider Organization 	Valley Clinic	RXA-10
 Immunization 		RXA segment
 Date administered 	July 23,1999	RXA-3
• Vaccine	MMR	RXA-5
CPT Code	90707	RXA-5
• Dose (ml)	0.5	RXA-6
Administering Provider Organization	Valley Clinic	RXA-10
• Client #3		PID segment
WIR Client ID	927389	PID-2
Chart Number	92HG9257	PID-3
• Name	JOSEPH FISHER	PID-5
Mother's maiden name	MARY LASOWSKI	PID-6
Birth date	May 28, 1998	PID-7
• Sex	M	PID-8
 Immunization 		RXA segment
Date administered	July 29, 1999	RXA-3
• Vaccine	MMR	RXA-5
CPT Code	90707	RXA-5
• Dose (ml)	0.5	RXA-6
Administering Provider Organization	Valley Clinic	RXA-10
Lot number	AD19487	RXA-15
Lot expiration date	December 12, 1999	RXA-16
Lot manufacturer	FLYBYNIGHT LABORATORIES (this manufacturer is not found in the valid list in HL7 Table 0227, and the invalid value will cause WIR to reject the message with an error message)	RXA-17

In an HL7 message, each segment is a single text line, ending with the carriage return character. In the examples, long lines are broken artificially for display purposes, and the carriage return character is denoted by <CR>.

```
FHS|^~\&|VALSYS|VALCLIN||WIR|19990802091523||filename1.h17|WEEKLY HL7
      UPLOAD | 00009972<CR>
BHS | ^~ \ & | VALSYS | VALCLIN | | WIR | 19990802091523 | | | | 00010223 < CR >
MSH|^~\&|VALSYS|VALCLIN||WIR|19990802091524||ADT^A31|00000123|P|2.3.1|||AL<CR>
PID|||45LR999||MILLER^GEORGE^M^JR|OLSON^MARTHA|19950227|M|||123 MAIN
      ST^^MADISON^WI^53000^US^^^DANE||||||000111222|||| |Y|2<CR>
PD1||||||||02^REMINDER/RECALL - ANY MENTOD^HL70215|Y| |A<CR>
NK1|1|MILLER^MARTHA|32^Mother^HL70063|123 MAIN ST^^MADISON^WI^53000^US^^^W1025
      (608)123-4567<CR>
NK1|2|MILLER^GEORGE|33^Father^HL70063<CR>
NK1|3|JONES^LUCAS^^^^|D3^Uncle^HL70063^^^|^Madison^WI^53715^USA^^^^|(515)829-
      1521||||||||||||||02<CR>
MSH|^~\&| VALSYS|VALCLIN||WIR|19990802091524||VXU^04|00000124|P|2.3.1|||ER<CR>
PID||66782|23LK729|CALIFANO^MARIA|DISTEFANO^ANGELICA|19980413|F<CR>
RXA|0|999|19990723|19990723|^^^90700^DtaP^CPT|0.5||||VALCLIN<CR>
RXA|0|999|19990723|19990723|^^^90707^MMR^CPT|0.5||||VALCLIN<CR>
MSH|^~\&|VALSYS|VALCLIN||WIR|19990802091526||VXU^04|00000125|P|2.3.1|||ER<CR>
```

```
PID||927389|92HG9257|FISHER^JOSEPH|LASOWSKI^MARY|19980528|M<CR>
RXA|0|999|19990729|19990729|^^^90707^MMR^CPT|0.5|||VALCLIN||||AD19487|
19991212|ZZ^FLYBYNIGHT LABORATORIES^HL70227||||A<CR>
BTS|3<CR>
FTS|1<CR>
```

Note: When a client is being introduced to WIR the VXU message must precede the ADT message as we must have one immunization for a client before they will be added to the database. Sending ADT messages is unnecessarily redundant when sending VXU messages for the same client, as the VXU message is capable of reporting the information for both message types.

In the example above, Valley Clinic sends a file of three HL7 messages to WIR. The messages are bracketed by file and batch header segments. The first message is of type ADT, which can be used when sending client demographic data without immunization information. This message type MUST follow a VXU message for the client if the client is new to the WIR system.

Client George M Miller Jr. is identified by Valley Clinic's chart number, 45LR999, in his PID segment. The message could have included George's WIR ID number in field PID-2, but does not have to, if it is not recorded in Valley Clinic's system. George's mother's maiden name, birth date, sex, address, and social security number also serve to identify him. Some other optional fields are not present, including some fields from the full HL7 standard not defined in this document because they are not used by WIR. Fields not present do not diminish the number of "|" delimiters, so later fields can be identified by ordinal position in the segment. Two NK1 segments give some information for George's mother and father, just the minimum required for his father, with address and telephone fields for his mother.

The next two PID segments in the second and third messages give a WIR client ID in field PID-2. This must have been transmitted earlier from WIR to Valley Clinic's system. In this case it is legitimate to omit more of the optional PID fields, since WIR must have at least the minimum required information for these clients even to create a record. However, if there is a possibility that Valley Clinic has new or changed information to send to WIR, these fields should be present, and it does no harm to repeat fields even if they have been transmitted previously.

```
FHS|^~\&|WIR|WIR||VALCLIN|19990803200106||filename2.h17||000023479|00009972<CR>
BHS|^~\&|WIR|WIR||VALCLIN|19990803200116||||00004321|00010223<CR>
MSH|^~\&|WIR|WIR||VALCLIN|19990803200117||ACK|00000456|P|2.3.1<CR>
MSA|AA|00000123<CR>
MSH|^~\&|WIR|WIR||VALCLIN|19990803200119||ACK|00000458|P|2.3.1<CR>
MSA|AE|00000125|INVALID MANUFACTURER CODE<CR>
ERR|RXA^152^17^1<CR>
BTS|2|<CR>
FTS|1<CR>
```

WIR answers the file from the above example with a file of ACK messages. Valley Clinic's message 00000123 had the value AL in field MSH-15, asking for acknowledgements of all messages. The value AA in MSA-1 tells that this message was processed without error. The next message, 00000124, uses the value ER to ask for acknowledgement only in case of errors, so this message is acknowledged implicitly by the absence of an ACK message for it. This example, while legitimate, is for purposes of illustration, and most providers will probably prefer to follow the WIR recommendation of error acknowledgements only. The last message, 00000125, did contain an error, and the ERR segment in its acknowledgement indicates the segment ID (RXA) of the segment, the line number (152) where it appears in the input file, the errant field (17) and the field component (1). The MSA segment contains the error message. Errors will be generated for missing required data, invalid data, or any other deviance from the form and content of messages as specified in this document. If all three messages in the first file above had requested error acknowledgement only, and none had any errors, the answering file from WIR would contain just the FSH, BHS, BTS, and FTS segments, and all messages would be implicitly acknowledged as successfully processed.

In the sample file exchange above, the outside system initiated the exchange with the file of ADT and VXU segments, and WIR responded with ACK segments. The format is identical when WIR sends ADT and VXU segments out, and the ACK responses are similar too. In the FHS, BHS, and MSH segments, the values of the fourth and sixth fields are reversed to show sender and receiver. WIR always sends its own client identifier in the required field PID-3, and includes the outside system's identifier in PID-2 if known. Outside systems are encouraged to store WIR's client ID, and use it in PID-2 when sending to WIR. This provides a firm basis for client identification, makes processing easier for the WIR system, and avoids errors in storing client information, such as creation of duplicate records when an insufficiently identified client record cannot be matched with a record already in the WIR database. Though WIR makes a great effort to match client records effectively, use of the WIR client ID is the best guarantee of clean and useful data.

Appendix A – HL7 Data Types

The Center for Disease Control Implementation Guide (CDC IG) contains clearly defined HL7 data types that are the building blocks of an HL7 message. This guide will avoid potentially ambiguous situations and will not redefine an already clearly defined section. Data types not otherwise noted herein, will adhere to corresponding definition in Chapter 4: HL7 Data Types of the CDC IG.

The following descriptions of HL7 data types are excerpted or adapted from the HL7 standard. See the field notes within each segment definition above on how to use data types in particular fields. Some data types have complex definitions much of which do not apply to WIR usage, and for these we omit much of the HL7 definition of the data type, referring instead to the field notes in the segment definitions.

CE

Coded Element

Example:

```
|F-11380^CREATININE^I9^2148-5^CREATININE^LN|
```

This data type transmits codes and the text associated with the code. To allow all six components of a CE data type to be valued, the maximum length of this data type must be at least 60.

Identifier (ST)

Sequence of characters (the code) that uniquely identifies the item being referenced by the <text>. Different coding schemes will have different elements here.

Text (ST)

Name or description of the item in question. E.g., myocardial infarction or X-ray impression. Its data type is string (ST).

Name of coding system (ST)

Each coding system is assigned a unique identifier. This component will serve to identify the coding scheme being used in the identifier component. The combination of the **identifier** and **name of coding system** components will be a unique code for a data item. Each system has a unique identifier. ASTM E1238-94, Diagnostic, procedure, observation, drug ID, and health outcomes coding systems are identified in the tables in Section 7.1.4 [of the full HL7 standard], "Coding schemes." Others may be added as needed. When an HL7 table is used for a CE data type, the *name of coding system* component is defined as *HL7nnnn* where *nnnn* is the HL7 table number.

Alternate components

These three components are defined analogously to the above for the alternate or local coding system. If the Alternate Text component is absent, and the Alternate Identifier is present, the Alternate Text will be taken to be the same as the Text component. If the Alternate Coding System component is absent, it will be taken to mean the locally defined system.

Note: The presence of two sets of equivalent codes in this data type is semantically different from a repetition of a CE-type field. With repetition, several distinct codes (with distinct meanings) may be transmitted.

Note: For HL7-defined tables which have not been adopted from some existing standard, the third component, "name of coding system," is constructed by appending the table number to the string "HL7." Thus, the field *RXR-2-site*, is a CE data type which refers to HL7 table number 0163. Its "name of coding system" component is "HL70163".

<u>CM</u>

Composite

```
Components: <point of care (IS) > ^ <room (IS) ^ <bed (IS) > ^ <facility (HD) ^ <location status (IS) ^ <patient location type (IS) > ^ <building (IS) > ^ <floor (IS) > ^ < street address (ST) > ^ <other designation (ST) > ^ <city (ST) > ^ <state or province (ST) > ^ <zip or postal code (ST) > ^ <country (ID) > ^ <address type (ID) > ^ <other geographic designation (ST) >

Subcomponents of facility (HD): <namespace ID (IS) > & <universal ID (ST) > & <universal ID type (ID) > ^
```

Example:

```
|^^^Valley Clinic|
```

Definition: The first component contains the inpatient or outpatient location at which the drug or treatment was administered (if applicable). The default (null) value is the current census location for the patient. Site-specific table. The first eight components have the same form as the first eight components of *PV1-3-assigned patient location*. The final eight components replace the ninth component of *PV1-3-assigned patient location* and represent the full address specification.

CX

Extended Composite ID with Check Digit

WIR uses this data type only for client identification in Patient Identification (PID) segments. See the field notes for values used for WIR.

HD

Hierarchic Designator

WIR uses this data type only to identify sender and receiver in Message Header (MSH) segments. See the field notes for values used for WIR.

ID

Coded Value for HL7 Defined Tables

The value of such a field follows the formatting rules for a ST field except that it is drawn from a table of legal values. There shall be an HL7 table number associated with ID data types. Examples of ID fields include religion and sex. This data type should be used only for HL7 tables. The reverse is not true, since in some circumstances it is more appropriate to use the CE data type for HL7 tables.

IS

Coded Value for User Defined Tables

The value of such a field follows the formatting rules for a ST field except that it is drawn from a site-defined (or user-defined) table of legal values. There shall be an HL7 table number associated with IS data types. An example of an IS field is the *Event reason code* defined in Section 3.3.1.4 [of the full HL7 standard], "Event reason code." This data type should be used only for user-defined tables. The reverse is not true, since in some circumstances, it is more appropriate to use the CE data type for user-defined tables.

NM

Numeric

A number represented as a series of ASCII numeric characters consisting of an optional leading sign (+ or -), the digits and an optional decimal point. In the absence of a sign, the number is assumed to be positive. If there is no decimal point the number is assumed to be an integer. Examples:

```
|999|
|-123.792|
```

Leading zeros, or trailing zeros after a decimal point, are not significant. For example, the following two values with different representations, "01.20" and "1.2", are identical. Except for the optional leading sign (+ or -) and the optional decimal point (.), no non-numeric ASCII characters are allowed. Thus, the value <12 should be encoded as a structured numeric (SN) (preferred) or as a string (ST) (allowed, but not preferred) data type.

SI

Sequence ID

A non-negative integer in the form of a NM field. See the field notes in segments using this data type for specifications of SI fields.

ST

String Data

String data is left justified with trailing blanks optional. Any displayable (printable) ACSII characters (hexadecimal values between 20 and 7E, inclusive, or ASCII decimal values between 32 and 126), except the defined delimiter characters. Example:

```
|almost any data at all|
```

To include any HL7 delimiter character (except the segment terminator) within a string data field, use the appropriate HL7 escape sequence.

Usage note: the ST data type is intended for short strings (e.g., less than 200 characters). For longer strings the TX or FT data types should be used.

<u>TS</u>

Time Stamp

```
Format: YYYY[MM[DD[HHMM[SS[.S[S[S]]]]]]]]]+/-ZZZZ]^<degree of precision>
```

Contains the exact time of an event, including the date and time. The date portion of a time stamp follows the rules of a date field and the time portion follows the rules of a time field. The specific data representations used in the HL7 encoding rules are compatible with ISO 8824-1987I.

In prior versions of HL7, an optional second component indicates the degree of precision of the time stamp (Y = year, L = month, D = day, H = hour, M = minute, S = second). This optional second component is retained only for purposes of backward compatibility.

By site-specific agreement, YYYYMMDD[HHMM[SS[.S[S[S]]]]]][+/-ZZZZ]^<degree of precision> may be used where backward compatibility must be maintained.

In the current and future versions of HL7, the precision is indicated by limiting the number of digits used, unless the optional second component is present. Thus, YYYY is used to specify a precision of "year," YYYYMM specifies a precision of "month," YYYYMMDD specifies a precision of "day," YYYYMMDDHH is used to specify a precision of "hour," YYYYMMDDHHMMSS is used to specify a precision of seconds, and YYYYMMDDHHMMSS.SSSS is used to specify a precision of ten thousandths of a second. In each of these cases, the time zone is an optional component. Maximum length of the time stamp is 26. Examples:

```
| 19760704010159-0600| 1:01:59 on July 4, 1976 in the Eastern Standard Time zone.

| 19760704010159-0500| 1:01:59 on July 4, 1976 in the Eastern Daylight Saving Time zone.

| 198807050000| Midnight of the night extending from July 4 to July 5, 1988 in the local time zone of the sender.

| 19880705| Same as prior example, but precision extends only to the day. Could be used for a birthdate, if the time of birth is unknown.
```

The HL7 Standard strongly recommends that all systems routinely send the time zone offset but does not require it. All HL7 systems are required to accept the time zone offset, but its implementation is application specific. For many applications the time of interest is the local time of the sender. For example, an application in the Eastern Standard Time zone receiving notification of an admission that takes place at 11:00 PM in San Francisco on December 11 would prefer to treat the admission as having occurred on December 11 rather than advancing the date to December 12.

One exception to this rule would be a clinical system that processed patient data collected in a clinic and a nearby hospital that happens to be in a different time zone. Such applications may choose to convert the data to a common representation. Similar concerns apply to the transitions to and from daylight saving time. HL7 supports such requirements by requiring that the time zone information be present when the information is sent. It does not, however, specify which of the treatments discussed here will be applied by the receiving system.

XAD

Address

```
Components: <street address (ST)> ^ <other designation (ST)> ^ <city (ST)> ^ <state or province (ST)> ^ <zip or postal code(ST)> ^ <country (ID)> ^ < address type (ID)> ^ <other geographic designation (ST)>^ <county/parish code (IS)> ^ <census tract (IS)> ^ <address representation code (ID)>
```

Example:

```
|1234 Easy St.^Ste. 123^San Francisco^CA^95123^USA^B^^SF^^|
```

Street address (ST)

The street or mailing address of a person or institution.

Other designation (ST)

Second line of address. In general, it qualifies address. Examples: Suite 555 or Fourth Floor.

City (ST)

State or province (ST)

State or province should be represented by the official postal service codes for that country.

Zip or postal code (ST)

Zip or postal codes should be represented by the official codes for that country. In the US, the zip code takes the form 99999[-9999], while the Canadian postal code takes the form A9A-9A9.

Country (ID)

Defines the country of the address. See Table 0212.

Address type (ID)

Address type is optional.

Other geographic designation (ST)

Other geographic designation includes country, bioregion, SMSA, etc.

County/parish code (IS)

A code that represents the county in which the specified address resides. Refer to *user-defined table 0289 – County/parish*. When this component is used to represent the county (or parish), component 8 "other geographic designation" should not duplicate it (i.e., the use of "other geographic designation" to represent the county is allowed only for the purpose of backward compatibility, and should be discouraged in this and future versions of HL7).

Census tract (IS)

An optional code that represents the census track in which the specified address resides. WIR does not store this value.

XCN

Extended Composite ID Number and Name For Persons

WIR uses this data type only to identify Provider Organizations that administer immunizations. See the field notes for segment RXA.

XPN

Extended Person Name

Example:

|Smith&St^John^J^III^DR^PHD^L|

Family name (ST)

Last Name Prefix (ST)

Given name (ST)

Middle initial or name (ST)

Suffix (ST)

Used to specify a name suffix (e.g., Jr. or III).

Prefix (ST)

Used to specify a name prefix (e.g., Dr.).

Degree (ST)

Used to specify an educational degree (e.g., MD).

Name type code (ID)

A code that represents the type of name. Refer to HL7 table 0200 - Name type for valid values.

Table 0200 - Name type

Value	Description
А	Alias Name
L	Legal Name
D	Display Name
M	Maiden Name
С	Adopted Name

Note: The legal name is the same as the current married name.

Name representation code (ID)

This component can be used when names are represented in ideographic or non-alphabetic systems. WIR ignores this component.

XTN

Extended Telecommunication Number

```
Components: [NNN] [(999)]999-9999 [X99999] [B99999] [C any text] ^ <telecommunication use code (ID)> ^ <telecommunication equipment type (ID)> ^ <email address (ST)> ^ <country code (NM)> ^ <area/city code (NM)> ^ <phone number (NM)> ^ <extension (NM)> ^ <any text (ST)>
```

Example:

(415)555-3210^ORN^FX^

[(999)] 999-9999 [X99999] [C any text]

Defined as the TN data type, except that the length of the country access code has been increased to three.

Telecommunication use code (ID)

A code that represents a specific use of a telecommunication number. Refer to HL7 table 0201 – Telecommunication use code for valid values.

Table 0201 - Telecommunication use code

Value	Description
PRN	Primary Residence Number
ORN	Other Residence Number
WPN	Work Number
VHN	Vacation Home Number
ASN	Answering Service Number
EMR	Emergency Number
NET	Network (email) Address
BPN	Beeper Number

Telecommunication equipment type (ID)

A code that represents the type of telecommunication equipment. Refer to HL7 table 0202 – Telecommunication equipment type for valid values.

Table 0202 – Telecommunication equipment type

Value	Description
PH	Telephone
FX	Fax
MD	Modem
СР	Cellular Phone
BP	Beeper
Internet	Internet Address: Use Only If Telecommunication Use Code Is NET
X.400	X.400 email address: Use Only If Telecommunication Use Code Is NET

Email address (ST)

Country code (NM) Area/city code (NM) Phone number (NM) Extension (NM) Any text (ST)

Appendix B – HL7 Tables

The following tables give valid values for fields in the segments defined above, in the cases where the field definitions reference an HL7 table number. The tables are considered to be part of the HL7 standard, but those tables designated as type User have values determined by WIR.

Туре	Table	Name	Value	Description
User	0001	Sex		
	0001		F	Female
	0001		M	Male
	0001		U	Unknown
HL7	0003	Event Type		- CHARLEST
/	0003	Dvoit Type	A31	ADT/ACK - Update patient information
	0003		V04	VXU – Unsolicited vaccination record update
11		D.	V 04	VAO – Orisolicited vaccination record update
User	0005	Race		Associated to Paragraph Alaska Nation
	0005		l .	American Indian or Alaska Native
	0005		A	Asian or Pacific Islander
	0005		В	Black
	0005		W	Caucasian
	0005		0	Other
	0005		U	Unknown
HL7	8000	Acknowledgment Code		
	8000		AA	Application Accept
	8000		AE	Application Error
	0008		AR	Application Reject
User	0063	Relationship		
	0063		18	Self
	0063		21	Unknown
	0063		26	Guardian
	0063		31	Court Appointed Guardian
	0063		32	Mother
	0063		33	Father
	0063		36	Emancipated Minor
	0063		48	Stepfather
	0063		49	
	0063		51	Stepmother Emergency Contact
				Emergency Contact
	0063		57	Adoptive Father
	0063		58	Adoptive Mother
	0063		61	Aunt
	0063		62	Brother
	0063		87	Foster Father
	0063		88	Foster Mother
	0063		97	Grandfather
	0063		98	Grandmother
	0063		A4	Half Brother
	0063		A5	Half Sister
	0063		B7	Sister
	0063		С3	Step Brother
	0063		C8	Step Sister
	0063		D3	Uncle
	0063		G7	Neighbor
	0063		G8	Other Relationship
	0063		G9	Other Relative
HL7	0003	Message Type		Outor Rolduvo
/	0076	micouge 1 ype	ACK	General acknowledgment message
	0076		ADT	ADT message
	0076		VXU	Unsolicited vaccination record update
HL7	0085	Observation result status codes		
	0085		0	Order detail description only
HL7	0103	Processing ID		
	0103		Р	Production
HL7	0104	Version ID		

Туре	Table	Name	Value	Description
.)	0104		2.3.1	Release 2.3.1 1999
HL7	0136	Yes/No Indicator		1.0.0000 2.0.11 1000
	0136	100/110 marcutor	Υ	Yes
	0136		N	No
HL7	0155	Accept/Application Acknowledgment Conditions		THO STATE OF THE S
	0155		ER	Error/reject conditions only
HL7	0162	Route of Administration		
	0162		ID	Intradermal
	0162		IM	Intramuscular
	0162		IN	Intranasal
	0162		IV	Intravenous
	0162		PO	Oral
	0162		sc	Subcutaneous
	0162		TD	Transdermal
	0162		MP	Multiple Puncture (Small Pox)
HL7	0163	Administrative Site		
	0163		LT	Left Thigh
	0163		LA	Left Arm
	0163		LD	Left Deltoid
	0163		LG	Left Gluteous Medius
	0163		LVL	Left Vastus Lateralis
	0163		LLFA	Left Lower Forearm
	0163		RA	Right Arm
	0163		RT	Right Thigh
	0163		RVL	Right Vastus Lateralis
	0163		RG	Right Gluteous Medius
	0163		RD	Right Deltoid
	0163		RLFA	Right Lower Forearm
User	0189	Ethnic Group	KEIA	rught Lower Folcann
0301	0189	Etimic Group	Н	Hispanic
	0189		NH	Non-Hispanic
User	0212	Nationality	INII	Non i lispanic
USEI	0212	ivationality	CA	Canada
	0212		US	United States of America
User	0215	Publicity Code	00	Officed States of Afficinea
USEI	0215	Fublicity Code	01	No reminder/recall
	0215		02	Yes reminder/recall – any method
HL7	0213	Identifier Type	02	res remindentecan – any method
1111	0203	Identifier Type	BR	Birth Registry Number
			MA	Medicaid Number
	0203		MC	Medicard Number Medicare Number
	0203			
	0203		MR PI	Medical Record Number
	0203			Patient Internal Identifier
	0203		PN	Person Number
	0203		PRN	Provider Number
	0203		PT	Patient External Identifier
	0203		RRI	Regional Registry ID
	0203		SR	State Registry Identifier
	0203	Manufacturers of vaccines (code =	SS	Social Security Number
HL7	0227	MVX)		
	0227		AB	Abbott
	0227		ACA	ACAMBIS
	0227		AD	Adams
	0227		ALP	Alpha

Type	Table	Name	Value	Description
	0227		AP	Sanofi Pastuer
	0227		AR	Armour (Inactive – use CSL)
	0227		AVB	Aventis Behring (Inactive – use CSL)
	0227		AVI	Aviron
	0227		ВА	Baxter (Inactive – use BAH)
	0227		ВАН	Baxter Health Care
	0227		BAY	Bayer
	0227		ВР	Berna (Inactive – use BPC)
	0227		BPC	Berna Products Corporation
	0227		BRR	Barr Laboratories
	0227		CEN	Centeon L.L.C. (Inactive – use CSL)
	0227		CHI	Chiron Corporation (Inactive – use NOV)
	0227		CMP	Celltech Medeva Pharm (Inactive – use NOV)
	0227		CNJ	Cangene Corporation
	0227		CON	Connaught (Inactive – use PMC)
	0227		CRU	Crucell
	0227		CSL	CSL Behring, Inc.
	0227		DVX	<u> </u>
				Dynavax Inc.
	0227		EVN	Evans (Inactive – use NOV)
	0227		GEO	GeoVax Labs, Inc
	0227		GRE	Greer
	0227		GRF	Grifols
	0227		IAG	Immuno International AG (Inactive – use BAH)
	0227		IDB	ID Biomedical
	0227		IM	Merieux (Inactive – Use PMC)
	0227		INT	Intercell Biomedical
	0227		IUS	Immuno-US
	0227		JPN	The Research foundation for Microbial Diseases of Osaka U.
	0227		KGC	Korea Green Cross
	0227		LED	Lederle (Inactive – use WAL)
	0227		MA	Massachusetts Public Health (Inactive-Use MBL)
	0227		MBL	Massachusetts Biologic Laboratories
	0227		MED	MedImmune
	0227		MIL	Miles (Inactive – use BAY)
	0227		MIP	Emergent BioDefense Operatons Lansing
	0227		MSD	Merck
	0227		NAB	North American Biologicals, Inc.
	0027		NAV	North American Vaccine (Inactive – use BAH)
	0227		NYB	New York Blood Center
	0227		NOV	Novartis
	0227		NVX	Novavax, Inc
	0227		OTC	Organon Teknika
	0227		ORT	Ortho
	0227		PAX	PaxVax
	0227		PD	Parkdale Pharmaceuticals (formerly Parke Davis)
	0227		PFR	Pfizer
	0227		PMC	Sanofi Pasteur Inc. (Connaught and Pasteur Merieux)
	0227		PRX	Praxis Biologics (Inactive – use WAL)
	0227		PSC	Protein Sciences
	0227		PWJ	Powderject Pharmaceutical
	0227		SCL	Sclavo
	0227		SEQ	Seqirus
	0227		SKB	SmithKline
	0227		SOL	Solvay Pharmaceuticals

Туре	Table	Name	Value	Description
	0227		SI	Swiss Serum and Vaccine Inst. (Inactive – use BPC)
	0227		TAL	Talecris Biotherapeutics (includes Bayer Biologicals)
	0227		USA	United States Army Medical Research
	0227		VAL	Valneva
	0227		VXG	Vacgen
	0227		WA	Wyeth-Ayerst (Inactive – use WAL)
	0227		WAL	Wyeth
	0227		ZLB	ZLB Behring (includes Aventis Behring and Armour
	0			Pharmaceutical Company (Inactive – use CSL)
	0227		ОТН	Other
	0227		UNK	Unknown manufacturer
Jser	0289	County/parish (Wisconsin only)		
	0289		WI001	Adams
	0289		WI003	Ashland
	0289		WI005	Barron
	0289		WI007	Bayfield
	0289		WI009	Brown
	0289		WI011	Buffalo
	0289		WI013	Burnett
	0289		WI015	Calumet
	0289		WI017	Chippewa
	0289		WI019	Clark
	0289		WI021	Columbia
	0289		WI021	Crawford
	0289		WI025	Dane
	0289		WI025	
				Dodge
	0289		WI029	Door
	0289		WI031	Douglas
	0289		WI033	Dunn
	0289		WI035	Eau Claire
	0289		WI037	Florence
	0289		WI039	Fond du Lac
	0289		WI041	Forest
	0289		WI043	Grant
	0289		WI045	Green
	0289		WI047	Green Lake
	0289		WI049	Iowa
	0289		WI051	Iron
	0289		WI053	Jackson
	0289		WI055	Jefferson
	0289		WI057	Juneau
	0289		WI059	Kenosha
	0289		WI061	Kewaunee
	0289		WI063	La Crosse
	0289		WI065	Lafayette
	0289		WI067	Langlade
	0289		WI069	Lincoln
	0289		WI071	Manitowoc
	0289		WI073	Marathon
	0289		WI075	Marinette
	0289		WI077	Marquette
	0289		WI078	Menominee
	0289		WI078	Milwaukee
	0289 0289		WI081 WI083	Monroe Oconto

Туре	Table	Name	Value	Description
	0289		WI085	Oneida
	0289		WI087	Outagamie
	0289		WI089	Ozaukee
	0289		WI091	Pepin
	0289		WI093	Pierce
	0289		WI095	Polk
	0289		WI097	Portage
	0289		WI099	Price
	0289		WI101	Racine
	0289		WI103	Richland
	0289		WI105	Rock
	0289		WI107	
				Rusk
	0289		WI109	St. Croix
	0289		WI111	Sauk
	0289		WI113	Sawyer
	0289		WI115	Shawano
	0289		WI117	Sheboygan
	0289		WI119	Taylor
	0289		WI121	Trempeleau
	0289		WI123	Vernon
	0289		WI125	Vilas
	0289		WI127	Walworth
	0289		WI129	Washburn
	0289		WI131	Washington
	0289		WI133	Waukesha
	0289		WI135	Waupaca
	0289		WI137	Waushara
	0289		WI139	Winnebago
	0289		WI141	Wood
NIP	NIP001	Immunization Information Source		
	NIP001		00	New Immunization Record
	NIP001		01	Historical Information
NIP	NIP002	Substance Refusal Reason		
	NIP002		00	Parental Refusal
	NIP002		01	Religious Exemption
NIP	NIP004	Contraindications, Precautions		Tronglede Exemplien
1411	NIP004	Contraindications, Freedutions	03	Allergy to baker's yeast (anaphylactic)
	NIP004		04	Allergy to egg ingestion (anaphylafctic)
	NIP004		05	Allergy to nfluen (anaphylafctic)
	NIP004		06	Allergy to neomycin (anaphylafctic)
	NIP004		07	Allergy to streptomycin (anaphylafctic)
	NIP004		08	Allergy to thimerosal (anaphylafctic)
	NIP004		09	Allergy to previous dose of this vaccine or to any of its
	NIP004		10	unlisted vaccine components (anaphylafctic) Anaphylactic (life-threatening) reaction of previous does
	NIP004		11	of this vaccine Collapse or shock like state within 48 hours of previous
	NIP004		12	dose of DTP/DtaP Convulsions (fits, seizures) within 3 days of previous dose of DTP/DtaP
	NIP004		13	Persistent, inconsolable crying lasting 3 hours within 48 hours of previous dose of DTP/DtaP
	NIP004		14	Current diarrhea, moderate to severe
	NIP004		15	Encephalopathy within 7 days of previous dose of DTP
	NIP004		16	Current fever with moderate-to-severe illness
	NIP004		17	Fever of 40.5 C (105 F) within 48 hours of previous dose of DTP/DtaP
	NIP004		18	Gullain-Barre syndrome (GBS) within 6 weeks of previous dose of DTP/DtaP

Type	Table	Name	Value	Description
Турс	NIP004	Traino -	21	Current acute illness, moderate to severe (with or without fever) (e.g. diarrhea, otitis media, vomiting)
	NIP004		22	Chronic illness
	NIP004		23	Immune globulin (IG) administration, recent or simultaneous
	NIP004		24	Immunity: diphtheria
	NIP004		25	Immunity: Haemophilus nfluenza type B (Hib)
	NIP004		XA	Immunity: hepatitis A
	NIP004		26	Immunity: hepatitis B
	NIP004		27	Immunity: measles
	NIP004		28	Immunity: mumps
	NIP004		29	Immunity: pertussis
	NIP004		30	Immunity: poliovirus
	NIP004		42	Immunity: rabies
	NIP004		31	Immunity: rubella
	NIP004		32	Immunity: tetanus
	NIP004		33	Immunity: varicella (chicken pox)
	NIP004		XC	History of Varicella
	NIP004		34	Immunodeficiency (family history)
	NIP004		35	Immunodeficiency (household contact)
	NIP004		36	Immunodeficiency
	NIP004		37	Neurologic disorders
	NIP004		38	Otitis media (ear infection) moderate to severe (with or without fever)
	NIP004		СР	Pertussis contraindication and precautions

Туре	Table	Name	Value	Description
	NIP004		39	Pregnancy (in recipient)
	NIP004		СТ	Tetanus contraindication – allergic reaction
	NIP004		40	Thrombocytopenia
	NIP004		41	Thrombocytopenic purpura (history)
	NIP004		CI	Contact with Infant(s) less than 6 months of age
	NIP004		HR	High Risk Condition(s)
NIP	NIP005	Event Consequence		
	NIP005	Zvent consequence	D	Patient Died
	NIP005		L	Life threatening illness
	NIP005		E	Required emergency room/doctor visit
	NIP005		Н	Required hospitalization
	NIP005		P	Resulted in prolongation of hospitalization
	NIP005		J	Resulted in permanent disability
NIP	NIP006	Patient Registry Status	J	Resulted in permanent disability
1411	NIP006	1 dient Registry Status	A	Active
	NIP006		N	Inactive
			P	
\\/\ID	NIP006	Pagation Cod	<u>r</u>	Permanently inactive
WIR	WIR001	Reaction Codes	LIVDOTON	Lhandonia humarezzanako asllaren 1985 40 km.
	WIR001		HYPOTON	Hypotonic-hyporesponsive collapse within 48 hours of immunization
	WIR001		SEIZURE	Seizure occurring within 3 days
	WIR001		CRYING	Persistent crying lasting >= 3 hours within 48 hours of
	77111001		OTT II TO	immunization
	WIR001		FEVER105	Temperature >= 105 (40.5 C) within 48 hours of immunization
WIR	99W01	WIR Student Information Codes		
	99W01		FERPA	FERPA Release
	99W01		GRADYEAR	Graduation Year
	99W01		ENROLLDATE	Date Enrolled in WI School
WIR	WVGC	Vaccine Group Code (WVGC)		
	WVGC	_	Adeno	Adeno
	WVGC		Anthrax	Anthrax
	WVGC		BCG	BCG
	WVGC		Cholera	Cholera
	WVGC		Diphtheria	Diphtheria Antitioxin
	WVGC		DTP/aP	Diphtheria, Tetanus, Acellular Pertussis
	WVGC		Encephalitis	Encephalitis
	WVGC		HepA	Hepatitis A
	WVGC		НерВ	Hepatitis B
	WVGC		Hib	Hib
	WVGC		HPV	Human Papilloma Virus
	WVGC		lg	Ig
	WVGC		IG-RSV	Respiratory syncytial virus Ig
	WVGC		Influenza	Influenza
	WVGC		Influenza A H1N1	Novel Influenza A H1N1
	WVGC		Lyme	
				Lyme Meacles Virus Vaccine
	WVGC		Measles	Measles Mymas Pyhalla
	WVGC		MMR	Measles, Mumps, Rubella
	WVGC		Meningo P	Meningitis Maningitis P
	WVGC		Meningo B	Meningitis B
	WVGC		Mumps	Mumps Virus Vaccine
	WVGC		Pertussis	Pertussis
	WVGC		Plague	Plague
	WVGC		Pneumococcal	Pneumococcal Conjugate
	WVGC		Pneumo-Poly	Pneumonia Polysaccharide
	WVGC		Polio	Poliomyelitis

Туре	Table	Name	Value	Description
	WVGC		Rabies	Rabies
	WVGC		Rotavirus	Rotavirus
	WVGC		Rubella	Rubella Virus Vaccine
	WVGC		Tetanus	Tetanus Diphtheria
	WVGC		Td	Tetanus Diphtheria
	WVGC		Typhoid	Typhoid
	WVGC		Smallpox	Vaccinia
	WVGC		Varicella	Varicella
	WVGC		Yellow Fever	Yellow Fever
	-			
77/ID	WVGC	Vaccine Trade Name (WVTN)	Zoster	Zoster
WIR	WVTN	vaccine Trade Name (w v Tiv)	ACAM2000	Smallpox
	WVTN		Acel-Imune	Diphtheria, tetanus, acellular pertussis
	WVTN		ActHib	Hemophilus influenza b PRP-T 4 dose
	WVTN		Adacel	TdaP > 7 years
	WVTN		Adeno T4	Adenovirus type 4, live oral
	WVTN		Adeno T7	
	WVTN		AGENO 17 AFLURIA	Adenovirus type 7, live oral
	WVTN			Influenza split virus
	WVTN		AFLURIA, P-free	Influenza preservative free
	WVTN		AFLURIA Quadrivalent	Influenza quadrivalen
	WVTN		AFLURIA Quad, P-Free	Influenza quadrivalent preservative free
	WVTN		Agriflu, P-free	Influenza preservative free
	WVTN		Anthrax	Anthrax
	WVTN		Attenuvax	Measles live
	WVTN		BabyBIG	Botulism Immune Globulin
	WVTN		BayTet	Tetanus Ig human
	WVTN		BCG-Cancer	Bacillus Calmette-Guerin bladder cancer
	WVTN		BCG-TB	Bacillus Calmette-Guerin TB
	WVTN		Bexsero	Meningococcal B, recombinant, OMV, adjuvanted
	WVTN		Biavax II	Rubella and mumps live
	WVTN		BIG	Botulism Immune Globulin
	WVTN		BioThrax	Anthrax
	WVTN		Boostrix	TdaP > 7 years
	WVTN		Botulinum-antitoxin	Botulinum antitoxin equine
	WVTN		Botulism	Botulism Immune Globulin
	WVTN		Certiva	Diphtheria, tetanus, acellular pertussis
	WVTN		Cervarix	Human Papilloma Virus, Bivalent
	WVTN		CMV-IgIV	Cytomegalovirus Ig IV human
	WVTN		Comvax	HepB-Hib Combination
	WVTN		DAPTACEL	Diphtheria, tetanus, acellular pertussis, 5 antigens
	WVTN		DECAVAC	Td , preservative free
	WVTN		Diphtheria	Diphtheria
	WVTN		Diphtheria-antitoxin	Diphtheria antitoxin, equine
	WVTN		Dryvax	Vaccinia(Smallpox) dry
	WVTN		DT	Diphtheria tetanus pediatric
	WVTN		DTP	Diphtheria, tetanus, whole cell pertussis
	WVTN		Engerix-B Adult	Hepatitis B adult dose 1ml
	WVTN		Engerix-B dialysis	HepB-Dialysis 4 dose
			Engerix-B Peds	Hepatitis B pediatric/adolescent .5ml
	WVTN		Flebogamma	Ig IV human
	WVTN		Flu-Imune	Influenza split virus
	WVTN		Flu-Shield	Influenza split virus
	WVTN		FLUAD	Influenza Trivalent Adjuvanted
	WVTN		Fluarix, P-free	Influenza preservative free
	WVTN			·
	WVTN		Fluarix Quadrivalent, P-	Influenza quadrivalent preservative free

Туре	Table	Name	Value	Description
			Free	

Туре	Table	Name	Value	Description
	WVTN		Flublok	Influenza recombinant preservative free
	WVTN		Flublok Quadrivalent	Influenza Quadrivalent Recombinant P-Free
	WVTN		Flucelvax	Influenza MDCK preservative free
	WVTN		Flucelvax Quadrivalent	Influenza, MDCK Quadrivalent
	WVTN		Flucelvax Quadrivalent P- Free	Influenza MDCK Quadrivalent preservative free
	WVTN		FluLaval	Influenza split virus
	WVTN		FluLaval, P-free	Influenza preservative free
	WVTN		FluLaval Quad, P-Free	Influenza quadrivalent preservative free
	WVTN		FluLaval Quadrivalent	Influenza, injectable, quadrivalent
	WVTN		FluMist	Influenza live, for intranasal use
	WVTN		FluMist Quadrivalent	Flu-nasal quadrivalent
	WVTN		Fluogen	Influenza split virus
	WVTN		Fluvirin	Influenza split virus
	WVTN		Fluvirin, P-free	Influenza preservative free
	WVTN		Fluzone	Influenza split virus
	WVTN		Fluzone High-Dose	Influenza split virus increased antigen content
	WVTN		Fluzone Intradermal	Influenza, seasonal, intradermal, p-free
	WVTN		Fluzone Intradermal Quad	influenza, intradermal, quadrivalent, preservative free
	WVTN		Fluzone, P-free	Influenza preservative free
	WVTN		Fluzone Quad	Fluzone Quadrivalent
	WVTN		Fluzone Quad PF 6-35M	Influenza quadrivalent, preservative free
	VVVIIN			6 month to 3 year dosage
	WVTN		Fluzone Quadrivalent, P- Free	Influenza quadrivalent preservative free
	WVTN		Gardasil	Human Papilloma Virus, Quadrivalent
	WVTN		Gardasil 9	Human Papilloma Virus, 9-valent
	WVTN		Havrix-Adult	Hepatitis A adult
	WVTN		Havrix-Peds 2 Dose	Hepatitis A pediatric/adolescent 2 dose
	WVTN		Havrix-Peds 3 Dose	Hepatitis A pediatric/adolescent 3 dose
	WVTN		HBIg	Hepatitis B Ig human
	WVTN		Heplisav-B	Hepatitis B, adjuvanted
	WVTN		Hib-TITER	Hemophilus influenza b HbOC 4 dose
	WVTN		Hiberix	Hemophilus influenza b PRP-T 4 dose
	WVTN		HyperTET	Tetanus immune globulin human
	WVTN		H1N1 MED Nasal	H1N1 live, for intranasal use
	WVTN		H1N1 P-free CSL	H1N1 monovalent inactivated preservative free
	WVTN		H1N1 P-free NOV	H1N1 monovalent inactivated preservative free
	WVTN		H1N1 P-free SAN	H1N1 monovalent inactivated preservative free
	WVTN		H1N1 CSL	H1N1 monovalent inactivated
	WVTN		H1N1 NOV	H1N1 monovalent inactivated
	WVTN		H1N1 SAN	H1N1 monovalent inactivated
	WVTN		Ig	Ig human
	WVTN		IgIV	Ig IV human
	WVTN		Imovax Rabies ID	Rabies intradermal
	WVTN		Imovax Rabies IM	Rabies intramuscular
	WVTN		Infanrix	Diphtheria, tetanus, acellular pertussis
	WVTN		IPOL	Poliovirus inactivated IPV
	WVTN		Ixiaro	Japanese Encephalitis for Intramuscular use
			JE-Vax	Japanese Encephalitis for Subcutaneous use
	WVTN		KINRIX	DTaP-IPV combination
	WVTN		LYMErix	Lyme disease
	WVTN		M-R-VAX	Measles and rubella live
	WVTN		Measles	Measles live 1964-1974
	WVTN		Measles-Rubella (MERU)	Measles and rubella live
	WVTN			

Туре	Table	Name	Value	Description
				1351 diphtheria toxoid conjugate vaccine

Туре	Table Name Value		Description		
	WVTN		MenHibrix	Meningococcal-Hib combination	
	WVTN		MENOMUNE	Meningococcal polysaccharide	
	WVTN		Menveo	Meningococcal oligosaccharide [groups A, C, Y and W-135] diphtheria toxoid conjugate vaccine	
	WVTN		Meruvax II	Rubella live	
	WVTN		MMR II	Measles, mumps and rubella live	
	WVTN		Mumps	Mumps	
	WVTN		Mumps-Rubella (MURU)	Rubella and mumps live	
	WVTN		Mumpsvax	Mumps live	
	WVTN		OmniHib	Hemophilus influenza b PRP-T 4 dose	
	WVTN		ORIMUNE	Poliovirus OPV live oral	
	WVTN		Pediarix	DTAP-HepB-Polio combination	
	WVTN		Pentacel	DtaP-Hib-IPV combination	
	WVTN		PedvaxHIB	Hemophilus influenza b OMP 3 dose	
	WVTN		Plague	Plague	
	WVTN		Pneumovax 23	Pneumococcal polysaccharide 23 valent	
	WVTN		PNU-IMUNE 23	Pneumococcal polysaccharide 23 valent	
	WVTN		Prevnar	Pneumococcal conjugate polyvalent	
	WVTN		Prevnar 13	Pneumococcal 13-valent conjugate	
	WVTN		ProHIBit	Hemophilus influenza b PRP-D booster	
	WVTN		ProQuad	Measles, mumps, rubella, varicella live	
	WVTN		Quadracel	DtaP-IPV combination	
	WVTN		RabAvert	Rabies intramuscular	
	WVTN		Recombivax Peds	Hepatitis B pediatric/adolescent .5ml	
	WVTN		Recombivax-Adult	Hepatitis B adult dose 1ml	
	WVTN		Recombivax-Dialysis	Hepatitis B Dialysis 4 dose	
	WVTN		Respigam	Respiratory syncytial virus Ig IV	
			Rho(D)Full	Rho(D)Ig RhIg human full-dose	
	WVTN		Rho(D)IV	Rho(D)Ig RhIg human IV	
	WVTN		Rho(D)Mini	Rho(D)Ig RhIg human mini-dose	
	WVTN		Rig	Rabies Ig human	
	WVTN		Rig-HT	Rabies Ig heat treated human	
			Rotarix	Rotavirus-RV1	
	WVTN		RotaShield	Rotavirus tetravalent live oral	
	WVTN		RotaTeq	Rotavirus pentavalent	
	WVTN		RSV-IgIV	Respiratory syncytial virus Iq IV	
	WVTN		Rubella	Rubella live	
	WVTN		Shingrix	Zoster (shingles), subunit	
	WVTN		Synagis	Respiratory syncytial virus Ig	
	WVTN		Td	Tetanus and diphtheria adult	
	WVTN		TENIVAC	Td , preservative free	
	WVTN		Tetramune	DTP – Hib combination	
	WVTN		Tig	Tetanus Ig human	
	WVTN		TriHIBit	DtaP-Hib combination	
	WVTN		Tripedia	Diphtheria, tetanus, acellular pertussis	
	WVTN		Trumenba	Meningococcal B, fully recombinant	
	WVTN		TT	Tetanus	
	WVTN		Twinrix	Hepatitis A & Hepatitis B adult	
	WVTN		Typhim Vi	Typoid VI capsular polysaccharide	
	WVTN		Typhoid	Typhoid vi capsular polysacchande Typhoid heat and phenol inactivated	
	WVTN		Typhoid-AKD	Typhoid acetone-killed, dried	
	WVTN		Vaccinia (smallpox),	Vaccinia (smallpox), diluted	
	WVTN		diluted	vaccinia (smanpox), unuteu	
	WVTN		Vaccinia immune globulin VIG	Vaccinia immune globulin VIG	
	WVTN		VAQTA-Adult	Hepatitis A adult	

Туре	Table	Name	Value	Description
	WVTN		VAQTA-Peds 2 Dose	Hepatitis A pediatric/adolescent 2 dose
	WVTN		Varivax	Varicella live
	WVTN		Vaxchora	Cholera, live attenuated
	WVTN		Vivotif Berna/Ty21a	Typhoid oral
	WVTN		VZIg	Varicella-zoster Ig human
	WVTN		YF-VAX	Yellow Fever live
	WVTN		Stamaril	Alternate yellow fever vaccine
	WVTN		Zostavax	Zoster (shingles), live

CPT Codes (WCPT) and CVX Codes (292)

CPT	CVX	Group	Vaccine	Trade Name	Description	MFG
90476	54	Adeno	Adeno T4	Adeno T4	Adenovirus type 4, live oral	WAL
90477	55		Adeno T7	Adeno T7	Adenovirus type 7, live oral	WAL
00504	82	A (I	Adeno, unspecified formulation	A in the many	Recorded as CVX 55	MID
90581	24	Anthrax	Anthrax	Anthrax BioThrax	Anthrax	MIP
90585	19	BCG	BCG-TB	BCG-TB	Bacillus Calmette-Guerin TB	OTC
	19	BCG				OTC
90586				BCG-Cancer	Bacillus Calmette-Guerin bladder cancer	OIC
90728		01 1	BCG		BCG	541/
90625	174	Cholera	Cholera, live attenuated	Vaxchora	Cholera, live attenuated	PAX
90725	26		Cholera, unspecified formulation		Cholera, unspecified formulation	
90719		Diphtheria	Diphtheria	Diphtheria	Diphtheria	PD
90700	20	DTP/aP	DTaP	Acel-Imune	Diphtheria, tetanus, acellular	WAL
				Certiva	pertussis	BAH
				Infanrix	_	SKB
			272	Tripedia		PMC
90701	01		DTP	DTP	Diphtheria, tetanus, whole cell pertussis	PMC
90702	28		DT	DT	Diphtheria tetanus pediatric	PMC
90720	22		DTP-Hib	Tetramune	DTP – Hib combination	WAL
90721	50		DTaP-Hib	TriHIBit	DtaP-Hib combination	PMC
90723	110		DTAP-HepB-Polio	Pediarix	DTAP-HepB-Polio combination	SKB
90696	130		DTaP-IPV	KINRIX	DTaP-IPV combination	SKB
				Quadracel		PMC
90698	120		DtaP-Hib-IPV	Pentacel	DtaP-Hib-IPV combination	PMC
	106		DTAP, 5 pertussis antigens	DAPTACEL	Diphtheria, tetanus, acellular pertussis, 5 antigens	PMC
	107		DTaP, unspecified formulation		Recorded as CVX 20	
90735	39	Encephalitis	Japanese Encephalitis-SC	JE-Vax	Japanese encephalitis for Subcutaneous use	JPN
90738	134		Japanese Encephalitis-IM	Ixiaro	Japanese encephalitis for Intramuscular use	VAL
	129		Japanese Enceph, unspecified formulation		Japanese Enceph, unspecified formulation	
90632	52	HepA	HepA adult	Havrix-Adult VAQTA-Adult	Hepatitis A adult	SKB MSD
90633	83		HepA-Ped 2 Dose		Hepatitis A pediatric/adolescent 2 dose	SKB MSD
90634	84		HepA -Peds	Havrix-Peds 3 Dose	Hepatitis A pediatric/adolescent 3 dose	SKB MSD
90636	104		HepA-HepB Adult	Twinrix	Hepatitis A & Hepatitis B adult	SKB
90730	85	1	Hep A, unspecified formulation		Hep A, unspecified formulation	
	31		Hep A-Peds, unspecified formulation		Recorded as CVX 85	
90636	104	HepB	HepA-HepB Adult	Twinrix	Hepatitis A & Hepatitis B adult	SKB
90723	110		DTAP-HepB-Polio	Pediarix	DTAP-HepB-Polio combination	SKB
90731	45	1	Hep B, unspecified formulation		Hep B, unspecified formulation	
90739	189		Hep B, adjuvanted	Heplisav-B	Hepatitis B, adult dosage (2 dose schedule), for intramuscular use	DVX
90740	44		Hep B-Dialysis 3 dose		Hepatitis B Dialysis 3 dose	
90743	43		HepB adult	Recombivax-Adult Engerix-B Adult	Hepatitis B adult dose 1ml	MSD SKB
90744	08		HepB pediatric	Recombivax Peds Engerix-B Peds	Hepatitis B pediatric/adolescent .5ml	MSD SKB
90745	42		Hep B, adolescent/high risk infant	gon/ 2 1 000	Hep B, adolescent/high risk infant	5.05
90746	43		HepB adult	Recombivax-Adult Engerix-B Adult	Hepatitis B adult dose 1ml	MSD SKB
90747	44		HepB-Dialysis 4 dose	Recombivax- Dialysis	Hepatitis B Dialysis 4 dose	MSD
				Engerix-B dialysis		SKB
90748	51		HepB-Hib HepB-Unspecified	Comvax	HepB-Hib Combination	MSD
90645	47	Hib	Hib-HbOC	Hib-TITER	Hemophilus influenza b HbOC 4	WAL
00040	77	1 110		I IID-TITEK	dose	**/\L

CPT	CVX	Group	Vaccine	Trade Name	Description	MFG
90646	46		Hib-PRP-D	ProHIBit	Hemophilus influenza b PRP-D	PMC
					booster	
90647	49		Hib-OMP	PedvaxHIB	Hemophilus influenza b OMP 3 dose	
90648	48		Hib-PRP-T	OmniHib	Hemophilus influenza b PRP-T 4	PMC
				ActHib	dose	PMC SKB
00700	22	-	DTP-Hib	Hiberix Tetramune	DTP – Hib combination	WAL
90720 90721	50	-	DTaP-Hib	TriHIBit	DtaP-Hib combination	PMC
90737	17	-	Hib, unspecified formulation	THIHDIL	Hib,unspecified formulation	FIVIC
90748	51	1	HepB-Hib	Comvax	HepB-Hib combination	MSD
90698	120		DtaP-Hib-IPV	Pentacel	DtaP-Hib-IPV combination	PMC
90644	148		Meningococcal C/Y-HIB PRP	MenHibrix	Meningococcal-Hib combination	SKB
90650	118	HPV	HPV, Bivalent	Cervarix	Human Papilloma Virus	SKB
90649	62		HPV, Quadrivalent	Gardasil	Human Papilloma Virus	MSD
90651	165		HPV, 9-valent	Gardasil 9	Human Papilloma Virus, 9-valent	MSD
	137		HPV, unspecified formulation		HPV, unspecified formulation	
90281	86	lg	Ig	Ig	lg human	
90283	87		IgIV	IgIV	Ig IV human	
		-		Flebogamma		
90287	27	-	Botulinum-antitoxin	Botulinum-antitoxin	Botulinum antitoxin equine	ļ
90288			Botulism	BabyBIG	Botulism Immune Globulin	
				Botulism	-	
90291	29	-	CMV-IgIV	BIG CMV-IgIV	Cytomegalovirus Ig IV human	-
90291	14	1	IG, unspecified formulation	OIVI V -IGI V	IG, unspecified formulation	
90296	12	1	Diphteria-antitoxin	Diphteria-antitoxin	Diphtheria antitoxin, equine	
90371	30	1	HBIg	HBIg	Hepatitis B Ig human	1
90375	34	1	RIg	Rig	Rabies Ig human	
90376	34		RIg-HT	RIg-HT	Rabies Ig heat treated human	1
90384	157		Rho(D)Full	Rho(D)Full	Rho(D)Ig Rhlg human full-dose	
90385	157		Rho(D)Mini	Rho(D)Mini	Rho(D)Ig Rhlg human mini-dose	
90386			Rho(D)IV	Rho(D)IV	Rho(D)Ig Rhlg human IV	
	156		Rho(D) IM or IV		Rho(D), unspecified formulation	
	159	_	Rho(D), unspecified formulation		Rho(D), unspecified formulation	
90389	13		TiG	BayTet	Tetanus Ig human	
				TiG	Tatanus immuus alahulia human	CDE
90393	79	-	Vaccinia immune globulin VIG	HyperTET Vaccinia-Ig	Tetanus immune globulin human Vaccinialg human	GRF
90393	36	-	VZIg	VZIg	Varicella-zoster Ig human	
90390	117	-	VZIG (IND)	VariZIG	Varicella-20ster ig flufflati	CNJ
		1	Varicella IG	Vanizio		0140
90378	93	IG-RSV	RSV-IgIM	Synagis	Respiratory syncytial virus Ig	
90379	71		RSV-IgIV	RSV-IgIV	Respiratory syncytial virus Ig IV	
			- 9	Respigam		
90630	166	Influenza	Influenza Intradermal Quadrivalent P-Free	Fluzone Intradermal Quad	influenza, intradermal, quadrivalent, preservative free	PMC
90653	168	1	Influenza Trivalent Adjuvanted	FLUAD	Influenza trivalent adjuvanted	SEQ
90654	144		Influenza Intradermal	Fluzone Intradermal	Influenza, seasonal, intradermal, p-free	PMC
90655	140	1	Influenza Preservative-Free	AFLURIA, P-free	Influenza preservative free	SEQ
				Agriflu, P-free	6 month to 3 year dosage	NOV
				Fluarix, P-free]	SKB
				Fluvirin, P-free		SEQ
]			Fluzone, P-free		PMC
90656				AFLURIA, P-free	Influenza preservative free	SEQ
				Agriflu, P-free	3 years and up dosage	NOV
				Fluarix, P-free	-	SKB
				FluLaval, P-free	-	SKB
				Fluvirin, P-free	-	SEQ
00657	1.11	-	Influenza	Fluzone, P-free	Influenza anlit virus	PMC
90657	141		Influenza	Flu-Imune	Influenza split virus 6 month to 3 year dosage	WAL WAL
				Flu-Shield Fluzone	o month to o year dosage	PMC
				AFLURIA	1	SEQ
				Fluvirin	1	SEQ
				Fluogen	1	PD
				FluLaval	1	SEQ
	ĺ			Flu-Imune	Influenza split virus	WAL
90658					J 30= a op * 11 40	
90658				Flu-Shield	3 years and up dosage	WAL
90658				Flu-Shield Fluzone	3 years and up dosage	WAL PMC

CPT	CVX	Group	Vaccine	Trade Name	Description	MFG
		Î		Fluvirin	•	SEQ
				Fluogen		PD
				FluLaval		SEQ
90659	16		Influenza-Whole Virus		Influenza whole virus	
90660	111		Flu-Nasal	FluMist	Influenza live, for intranasal use	MED
90661	153		Influenza MDCK Preservative- Free	Flucelvax	Influenza, injectable, MDCK, preservative free	NOV
90662	135		Influenza High Dose	Fluzone High-Dose	Influenza split virus increased antigen content	PMC
90672	149	=	Flu-Nasal Quadrivalent	FluMist Quadrivalent	Influenza quadrivalent live, for intranasal use	MED
90673	155	=	Influenza Recombinant P-Free	Flublok	Influenza, recombinant, injectable, preservative free	PSC
90674	171	-	Influenza MDCK Quadrivalent P-Free	Flucelvax Quadrivalent, P- Free	Influenza MDCK quadrivalent preservative free	SEQ
90682	185		Influenza Quad Recombinant P-Free	Flublok Quadrivalent	Influenza Quadrivalent recombinant P-Free	PSC
90685	161		Influenza Quadrivalent P-Free 6-35M	Fluzone Quad PF 6- 35M	preservative free 6 month to 3 year dosage	PMC
90686	150		Influenza Quadrivalent P-Free	AFLURIA Quad, P- Free	Influenza, injectable, quadrivalent, preservative free	SEQ
				Fluarix Quadrivalent, P- Free	3 years and up dosage	SKB
				FluLaval Quad, P- Free		IDB
				Fluzone Quadrivalent, P- Free		PMC
90687	158		Influenza Quadrivalent	Fluzone Quad	Influenza virus vaccine, quadrivalent, split virus, when administered to individuals 6-35 months of age, for intramuscular use	PMC
90688				AFLURIA Quadrivalent	Influenza virus vaccine, quadrivalent, split virus, when administered to	SEQ
				FluLaval Quadrivalent	individuals 3+ years of age, for intramuscular use	IDB
				Fluzone Quad	1	PMC
90724	88	-	Influenza, unspecified formulation		Influenza, unspecified formulation	
	151		Influenza Nasal, unspecified formulation		Influenza Nasal, unspecified formulation	
90756	186		Influenza MDCK Quadrivalent	Flucelvax Quadrivalent	Influenza, MDCK, Quadrivalent	SEQ
90664	125	Influenza A	Novel Influenza A H1N1-Nasal	H1N1 MED Nasal	H1N1 live, for intranasal use	MED
90666	126	H1N1	Novel Influenza A H1N1, P-free		H1N1 monovalent inactivated	CSL
	1			H1N1 P-free NOV	preservative free	NOV
				H1N1 P-free SAN	1	PMC
		1		H1N1 CSL	H1N1 monovalent inactivated	CSL
90668	127		Novel Influenza A H1N1	H1N1 NOV	1	NOV
				H1N1 SAN	1	PMC
90663	128	1	Novel Influenza A H1N1 all formulations	-	H1N1 all formulations	
90665	66	Lyme	Lyme	LYMErix	Lyme disease	SKB
90705	05	Measles	Measles	Measles	Measles live 1964-1974 (Eli Lilly)	MSD
00700	0.4	4	Manalan Dubatta	Attenuvax	Measles live	MSD
90708	04		Measles-Rubella	M-R-VAX Measles-Rubella	Measles and rubella live	MSD MSD
				(MERU)		
90704	07	Mumps	Mumps	Mumps	Mumps 1950-1978	MSD
00700		4	Puballa Mumpa NOS	Mumpsvax	Mumps live	MSD
90709	20	-	Rubella-Mumps, NOS	Piovov II	Pubella and mumpa live	Med
	38		Rubella-Mumps	Biavax II	Rubella and mumps live	MSD
				Mumps-Rubella (MURU)		MSD
90707	03	MMR	MMR	MMR II	Measles, mumps and rubella live	MSD
90710	94	1	MMRV	ProQuad	Measles, mumps, rubella, varicella	MSD
551 10	l 54		TAUAIL Z A	1000000	•	
					live	

CPT	CVX			Trade Name	Description	MFG
90734	114		Meningococcal-MCV4P	Menactra	Meningococcal polysaccharide	PMC
					[groups A, C, Y and W-135]	
	400		Maning and and I MCV/4C	Manua	diphtheria toxoid conjugate vaccine	NOV
	136		Meningococcal-MCV4O	Menveo	Meningococcal oligosaccharide [groups A, C, Y and W-135]	NOV
					diphtheria toxoid conjugate vaccine	
	147	1	Meningococcal-MCV4		MCV4, unspecified formulation	
			3		[groups A, C, Y and W-135]	
	108		Meningococcal, unspecified		Meningococcal, unspecified	
			formulation		formulation	
90644	148		Meningococcal C/Y-HIB PRP	MenHibrix	Meningococcal-Hib combination	SKB
90621	162	Meningo B	Meningococcal B, recombinant	Trumenba	Meningococcal B, fully recombinant	PFR
90620	163		Meningococcal B, OMV	Bexsero	Meningococcal B, recombinant,	SKB
	164	1	Meningococcal B, unspecified		OMV, adjuvanted Meningococcal B, unspecified	
	104		formulation		formulation	
90715	115	Pertussis	TdaP > 7 Years	Adacel	TdaP > 7 years	PMC
				Boostrix	1	SKB
	11	1	Pertussis		Pertussis vaccine	
90712	02	Polio	Polio oral	ORIMUNE	Poliovirus OPV live oral	WAL
90713	10	1	Polio injectable	IPOL	Poliovirus inactivated IPV	PMC
90723	110		DTAP-HepB-Polio	Pediarix	DTAP-HepB-Polio combination	SKB
90696	130]	DTaP-IPV	KINRIX	DTaP-IPV	SKB
				Quadracel		PMC
90698	120		DtaP-Hib-IPV	Pentacel	DtaP-Hib-IPV combination	PMC
	89		Polio, unspecified formulation		Polio, unspecified formulation	
90727	23	Plague	Plague	Plague	Plague	GRE
90732	33	Pneumo-Poly	Pneumococcal 23	PNU-IMUNE 23	Pneumococcal polysaccharide 23	WAL
				Pneumovax 23	valent	MSD
90669	100	Pneumococcal	Pneumo-Conjugate 7	Prevnar	Pneumococcal conjugate polyvalent	WAL
90670	133	1 neamococcai	Pneumo-Conjugate 13	Prevnar 13	Pneumococcal 13-valent conjugate	PFR
000.0	109	1	Pneumococcal, unspecified	1 TOVIIGI TO	Pneumococcal, unspecified	
			formulation		formulation	
	152		Pneumococcal Conjugate,		Pneumococcal Conjugate,	
			unspecified		unspecified formulation	
90675	18	Rabies	Rabies-intramuscular		Rabies intramuscular	
	175		Rabies-intramuscular, Diploid	Imovax Rabies IM	Rabies intramuscular, diploid cell	PMC
			cell culture Rabies-intramuscular,	RabAvert	culture Rabies intramuscular, Fibroblast	SKB
	176		Fibroblast culture	Rapaveri	culture	SND
90676	40		Rabies-intradermal	Imovax Rabies ID	Rabies intradermal	PMC
90726	90		Rabies, unspecified formulation		Rabies, unspecified formulation	
90680	74	Rotavirus	Rotavirus, Tet	RotaShield	Rotavirus tetravalent live oral	WAL
					(removed on 10/16/1999)	
	116		Rotavirus, Pent	RotaTeq	Rotavirus pentavalent (after	MSD
					02/02/2006)	
	122		Rotavirus, unspecified		(between 10/16/1999 and	
00694	110	-	formulation Potovirus monovelent	DOTABLY	02/01/2006)	SKB
90681 90706	119	Puballa	Rotavirus, monovalent	ROTARIX	Pubella livo	
90706	06	Rubella	Rubella	Rubella Meruvax II	Rubella live	MSD MSD
90708	04	1	Measles-Rubella	Measles-Rubella	Measles and rubella live	MSD
30100	04		Ividasids-i/ubdild	(MERU)	INICASICS AND TUDENA IIVE	טטואו
				M-R-VAX	1	MSD
90709		1	Rubella-Mumps NOS		Rubella-Mumps, NOS	
	38	1	Rubella-Mumps	Mumps-Rubella	Rubella and mumps live	MSD
			1	(MURU)	, .	
				Biavax II		MSD
	75	Smallpox	Smallpox	ACAM2000	Smallpox	PMC
		1	Smallpox	Dryvax	Vaccinia(Smallpox) dry	WAL
	105		Vaccinia (Smallpox), diluted	Vaccinia (smallpox),	Vaccinia (smallpox), diluted	
00740	00	T-4	Ta	diluted	Totonuo and dinhthada adult	DMC
90718	09	Td	Td	Td	Tetanus and diphtheria adult	PMC MBL
90714	113	4	Td Preservative-Free	DECAVAC	Td preservative free – CPT code is	PMC
30/14	113		i u Fieseivälive-Fiee		effective for immunizations given on	FIVIC
				TENIVAC	or after 7/1/2005	
00745	445	4	TdoD - 7 Vacas	Td P-free		DMC
90715	115		TdaP > 7 Years	Adacel	TdaP > 7 years	PMC
	138	-	Td (adult) not adsorbed	Boostrix	Td (adult) not adsorbed	SKB
	130	ı	Ta (addit) HOL adsorbed	1	Tra (addit) flot adsorbed	<u> </u>

CPT	CVX	Group	Vaccine	Trade Name	Description	MFG
	139		Td (adult) unspecified formulation		Td (adult) unspecified formulation	
90703	35	Tetanus	Tetanus	TT	Tetanus	PMC
	142		Tetanus toxoid, not adsorbed		Tetanus toxoid, not adsorbed	
	112		Tetanus toxoid, unspecified formulation			
90690	25	Typhoid	Typhoid-oral	Vivotif Berna/Ty21a	Typhoid oral	
90691	101		Typhoid-ViCPs	Typhim Vi	Typoid VI capsular polysaccharide	PMC
90692	41		Typhoid-HP	Typhoid	Typhoid heat and phenol inactivated	
90693	53		Typhoid-AKD	Typhoid-AKD	Typhoid acetone-killed, dried (military)	
90714	91		Typhoid, unspecified formulation		Typhoid, unspecified formulation (after 7/1/2005, no CPT code is associated with this vaccine group)	
90710	94	Varicella	MMRV	ProQuad	J 1,	MSD
90716	21	1	Varicella	Varivax	Varicella live	MSD
	37	Yellow Fever	Yellow Fever US	YF-VAX	Yellow Fever live	PMC
90717	183		Yellow fever - alt	Stamaril	Alternate yellow fever vaccine	PMC
			Yellow fever		Yellow fever US or yellow fever alternate	
	184	1	Yellow fever, unspecified formulation		Yellow fever, unspecified formulation	
90736	121	Zoster	Zoster (shingles), live	Zostavax	Zoster (shingles), live	MSD
90750	187	1	Zoster (shingles), subunit	Shingrix	Zoster (shingles), subunit	SKB
	188	1	Zoster, unspecified formulation		Zoster, unspecified formulation	

Appendix C – Error Messages

Error

The following is a list of common error messages that WIR will return for validation of message format, datum values, and business rules.

	Msg.				Sub		
Msg. Type	Code	Error Status Text	Segment	Comp.	Comp.		Error Message
Update/Query			MSH			Hard	NUMBER OF MESSAGES RECEIVED EXCEEDS 1
Update/Query			MSH			Hard	LONE MSH SEGMENT IN FILE
Update/Query			MSH	01		Hard	
Update/Query	102	Invalid Data Value	MSH	02		Hard	MESSAGE REJECTED - INVALID ENCODING CHARACTERS
Update/Query	101	Required Field Missing	MSH	04	02	Hard	MESSAGE REJECTED - INVALID OWNING PROVIDER ORGANIZATION ID
Update/Query			MSH	04		Hard	Record rejected. The provider organization that initiated this data exchange is not identified as a parent or vendor of the organization that it labeled as the "SENDING PROVIDER ORGANIZATION" for this record.
Update/Query			MSH	04		Hard	Message rejected The initiating and owning providers do not have a relationship in the IR.
Update/Query	100	Segment Sequence Error	MSH	09		Hard	MESSAGE REJECTED - INVALID MESSAGE TYPE SPECIFIED
Update/Query	101	Required Field Missing	MSH	10		Hard	MESSAGE REJECTED - MESSAGE CONTROL ID IS A REQUIRED FIELD
Update/Query			MSH	12		Hard	UNSUPPORTED HL7 VERSION
Update	102	Invalid Data Value	PID	03	05	Hard	MESSAGE REJECTED - PATIENT IDENTIFIER TYPE OF PI OR PN OR PRN OR PT REQUIRED
Update	101	Required Field Missing	PID	03		Hard	MESSAGE REJECTED - PATIENT IDENTIFIER LIST REQUIRED
Update	101	Required Field Missing	PID	05	01	Hard	MESSAGE REJECTED - PATIENT LAST NAME REQUIRED
Update	102	Invalid Data Value	PID	05	01	Hard	Message rejected. Client last name must be greater than one character in length.
Update	102	Invalid Data Value	PID	05	01	Hard	Message rejected. BABY is not a valid last name.
Update	101	Required Field Missing	PID	05	02	Hard	MESSAGE REJECTED - PATIENT FIRST NAME REQUIRED.
Update	102	Invalid Data Value	PID	05	02	Hard	Message rejected. Client first name must be greater than one character in length.
Update	102	Invalid Data Value	PID	05	02	Hard	Message rejected. BABY is not a valid first name.
Update			PID	05	02	Hard	Record Rejected - Invalid first name (MALE1 MELISSA).
Update	101	Required Field Missing	PID	07		Hard	MESSAGE REJECTED - Date of birth is a required field
Update	102	Invalid Data Value	PID	07		Hard	MESSAGE REJECTED - INVALID DATE OF BIRTH. MUST BE PRIOR TO OR EQUAL TO TODAY.
Update	102	Invalid Data Value	PID	07		Hard	MESSAGE REJECTED - Invalid date of birth format
Update	102	Invalid Data Value	PID	07		Hard	MESSAGE REJECTED - A VALID DATE OF BIRTH MUST BE SPECIFIED.
Update	102	Invalid Data Value	PID	11	04	Soft	Informational error - Invalid state code (Wisconsin). No value stored.

	Error Msg.				Sub		
Msg. Type	Code	Error Status Text	Segment	Comp.	Comp.		Error Message
Update	102	Invalid Data Value	PID	19		Soft	Informational error - Duplicate SSN. No value stored.
Update	102	Invalid Data Value	PID	19		Soft	INFORMATIONAL ERROR - Invalid SSN. SSN either starts with 000 or ends with 0000.
Update	102	Invalid Data Value	PID	19		Soft	INFORMATIONAL ERROR - Invalid SSN. SSN has 9 identical numbers.
Update	102	Invalid Data Value	PID	19		Soft	INFORMATIONAL ERROR - Invalid SSN. SSN has an invalid pattern.
Update	102	Invalid Data Value	PID	19		Soft	Invalid SSN. SSN has non-numeric characters.
Update	102	Invalid Data Value	PID	19		Soft	Invalid SSN. SSN not 9 characters in length.
Update			PD1				
Update	101	Required Field Missing	NK1	02	02	Soft	RELATIONSHIP MISSING FIRST NAME. NO VALUE STORED.
Update	102	Invalid Data Value	NK1	03	01	Soft	INFORMATIONAL ERROR - NO RELATIONSHIP CODE SPECIFIED. DEFAULTING TO GUARDIAN
Update	102	Invalid Data Value	NK1	03	01	Soft	INFORMATIONAL ERROR - INVALID RELATIONSHIP CODE. DEFAULTING TO GUARDIAN.
Update			RXA			Hard	MESSAGE REJECTED - ALL RXA SEGMENTS INVALID.
Update			RXA				The incoming delete immunization does not match an existing immunization in WIR. This delete was not processed.
Update			RXA				The sending provider organization does not own the existing matched immunization in WIR. This delete was not processed.
Update			RXA			Hard	MESSAGE REJECTED - RXA SEGMENT REQUIRED FOR VXU MESSAGE TYPE.
Update	102	Invalid Data Value	RXA	05		Hard	Invalid immunization INVALID ADMINISTERED CODE.
Update	101	Required Field Missing	RXA	06		Hard	ADMINISTERED AMOUNT IS A REQUIRED FIELD.
Update	102	Invalid Data Value	RXA	06		Hard	INFORMATIONAL ERROR - Invalid immunization INVALID ADMINISTERED AMOUNT
Update			RXA	09		Hard	RECORD REJECTED - 07 is not a valid immunization source for this provider organization.
Update	101	Required Field Missing	RXA	10	02	Soft	Administering provider last name is required to use administering provider field.
Update	102	Invalid Data Value	RXA	10	02	Soft	Informational error - Invalid administered by last name (Davis33 (Cerner)). No value stored.
Update			RXA	10	02	Soft	Informational error - More than one clinician found to match (LAST_NAME, FIRST_NAME)
Update			RXA	1 <i>7</i>		Soft	Informational error - Trade Name (Pneumovax 23) not produced by manufacturer (WAL). Defaulting to unknown manufacturer.
Update			RXR				
Update	102	Data type error	ОВХ			Soft	INVALID OBX SEGMENT - CONTRAINDICATION/PRECAUTION LOINC CODE SPECIFIED WITH IMMUNITY OBSERVATION VALUE. NO VALUE STORED.
Update	102	Data type error	ОВХ	03		Hard	INVALID OBX SEGMENT - Required OBX-03 LOINC code is null or invalid
Update	101	Required Field Missing	ОВХ	05		Hard	INVALID OBX SEGMENT - OBX-05 Observation value does NOT match observation coding system.

	Error Msg.				Sub		
Msg. Type	Code	Error Status Text	Segment	Comp.	Comp.	•	Error Message
Update	101	Required Field Missing	ОВХ	11		Hard	INVALID OBX SEGMENT - OBX-11 Observation Result status is a required field.
Update	102	Invalid Data Value	ОВХ			Soft	INACCURATE OR MISSING OBSERVATION VALUE. NO VALUE STORED.
Update						Hard	Record rejected. Client may not be updated since the existing client that it matches does not consent to share immunizations with your organization.
Update							PID SEGMENT - INVALID SOCIAL SECURITY NUMBER.
Update							Record rejected. This immunization matches another immunization in incoming file. The incoming immunization that this system retained may be identified by the following characteristics -> Vaccination Date: 02232012. 0
Query			QRD			Soft	Client has an 'Allow Sharing of Immunization Data' indicator = No.
Query	101	Required Field Missing	QRD	01		Hard	MESSAGE REJECTED - Query Date is a required field
Query	102	Invalid Data Value	QRD	01		Hard	MESSAGE REJECTED - Invalid Date format
Query	102	Invalid Data Value	QRD	01		Hard	MESSAGE REJECTED - Invalid Query Date
Query	101	Required Field Missing	QRD	02		Hard	MESSAGE REJECTED - Query Format Code is a required field
Query	102	Invalid Data Value	QRD	02		Hard	MESSAGE REJECTED - Invalid Query Format Code
Query	101	Required Field Missing	QRD	03		Hard	MESSAGE REJECTED - Query Priority is a required field
Query	102	Invalid Data Value	QRD	03		Hard	MESSAGE REJECTED - Invalid Query Priority Code
Query	101	Required Field Missing	QRD	04		Hard	MESSAGE REJECTED - Query ID is a required field
Query	101	Required Field Missing	QRD	07	01	Hard	MESSAGE REJECTED - Quantity Limited Request is a required field
Query	102	Invalid Data Value	QRD	07	01	Hard	MESSAGE REJECTED - Invalid Query Quantity
Query	102	Invalid Data Value	QRD	07	02	Hard	MESSAGE REJECTED - Invalid Query Units
Query	101	Required Field Missing	QRD	08	02	Hard	MESSAGE REJECTED - Last name required for Who Subject Filter
Query	101	Required Field Missing	QRD	08	03	Hard	MESSAGE REJECTED - First name required for Who Subject Filter
Query	101	Required Field Missing	QRD	08		Hard	MESSAGE REJECTED - Who Subject Filter is a required field.
Query	101	Required Field Missing	QRD	09	01	Hard	MESSAGE REJECTED - What Subject Filter is a required field
Query	102	Invalid Data Value	QRD	09	01	Hard	MESSAGE REJECTED - Invalid What Subject Filter Identifier(s)
Query	102	Invalid Data Value	QRD	10	01	Hard	MESSAGE REJECTED - Invalid What Department Data Code(s).
Query	101	Required Field Missing	QRD	10		Hard	MESSAGE REJECTED - What Department Data Code is a required field.
Query	100	Segment Sequence Error	QRF			Hard	MESSAGE REJECTED - QRF SEGMENT BEFORE QRD SEGMENT
Query	101	Required Field Missing	QRF	01		Hard	MESSAGE REJECTED - WHERE SUBJECT FILTER IS A REQUIRED FIELD.
Query	101	Required Field Missing	QRF	05	02	Hard	MESSAGE REJECTED - Date of birth is a required field
Query	102	Invalid Data Value	QRF	05	02	Hard	MESSAGE REJECTED - Invalid date of birth format

Document Updates

Version	Version	Revised By	Description
No.	Date		
1.0	1-Sep-2016	Amanda Ray	Updated Vaccine and Manufacturer Tables
1.1	12-Dec-2016	Amanda Ray	Updated Vaccine and Manufacturer Tables
1.2	20-Mar-2017	Amanda Ray	Added Afluria Quad, Afluria Quad P-Free, and Quadracel vaccines
1.3	11-Aug-2017	Jayme Judd	Added Flublok Quadrivalent and Flucelvax Quadrivalent. Updated RabAvert and Imovax Rabies IM.
1.4	13-Sept-2017	Jayme Judd	Added Yellow Fever vaccines
1.5	20-Sept-2017	Jayme Judd	Updated Vaccine and Manufacturer Tables
1.6	10-Nov-2017	Jayme Judd	Updated Vaccine and Manufacturer Tables
1.7	08-Dec-2017	Rebekah Van Dusen	Added Zoster vaccines (Shingrix and unspec form). Added missing manufacturers.
1.8	20-Feb-2018	Amanda Ray	Updated Flulaval, P-free typo
1.9	27-Feb-2018	Mark Ehlke	Added Vaxchora information
2.0	23-Mar-2018	Mark Ehlke	Updated table for HepB related CVX codes, WVTN and related CVX. Updated manufacturer table.
2.1	05-May-2018	Mark Ehlke	Updated MFG for Bexsero to SKB.