## PharmacyBrands Canada









PHARMACIST ASSESSMENT Pregnant Patients	Store Name:
DOCUMENTATION & NOTIFICATION Public TDAP Vaccine	Address/Phone:
PATIENT INFOMATION	
Name: DOB:	Date: Rx/Tx#:
PHN:	RATIONAL FOR ASSESSMENT (additional pg
Pregnant 🗆 Week: Lactating 🗆 Age of Infant:	□ Patient is Pregnant
PHARMACIST INFORMATION	Gestation week:(for reference see Alberta Blue Cross Pharmacy Benefact 769)
Name:	□ Reviewed patient's immunization Hx on NetCare
Store Address:	□Patient has not received Tdap vaccine earlier in
Phone:	this pregnancy
Fax:	□ Patient is not suffering from any severe acute illness
License #:	Patient has no history of anaphylaxis or other
Signature:	severe reaction after previous administration of
Signature.	Tdap vaccine.
	□ Patient has no allergy or hypersensitivity to any component of the vaccine
ORIGINAL PRESCRIBER INFORMATION	<u>Adacel:</u> tetanus toxoid, diptheria toxoid,
Dr.	acellular pertussis toxoid (PT), filamentous
ASSESSMENT TYPE	haemaggulutinin (FHA), pertactin (ORN), fimbriae types 2, 3, AIPO4, 2-phenoxyrthanol,
Initiating Medication Therapy	formaldehyde, and glutaraldehyde in trace
ORIGINAL PRESCRIPTION	<ul> <li>amounts.</li> <li><u>Boosterix:</u> diptheria toxoid, 3 purified pertussis</li> </ul>
Not applicable - initiating medication therapy	antigens [pertussis toxoid (PT), filamentous
MODIFIED OR NEW PRESCRIPTION LABEL	haemagglutinin (FHA) and pertactinand tetanus
	toxoid, Al, NaCl, water for injection, Na2PO4, formaldehyde, glutaraldehyde, glycine,
Pharmacist affix modified or new prescription label	monopotassium phosphate, polysorbate 90 and
here (Adaptation/Renewal/Medication Related	KCI. REDUCE PRESCRIPTION TO WRITING
Emergency/Initiating Medication Therapy/Trial Rx)	(N/A for refusal to fill / trial Rx / Opioid management)
	□ Not applicable (Patient <b>NOT</b> eligible for
	public program)
	□ Adacel vaccine, publicly funded, M: 1 dose
	SIG: administer IM
	☐ Boostrix vaccine, publicly funded, M: 1 dose SIG: administer IM
	FOLLOW-UP & MONITORING PLAN
INFORMED CONSENT	
The Patient and/or their representative (name):	
was provided with sufficient information, including the	
risks and benefits associated with the assessment and	
voluntarily provided their consent.	

Notification Information: A Fax A Phone DATE: NOTIFIED (other than above)

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