

SPECIAL AUTHORITY REQUEST SACUBITRIL-VALSARTAN COMBINATION

HLTH 5484 2022/04/29

For up-to-date criteria and forms, please check: www.gov.bc.ca/pharmacarespecialauthority

Fax requests to 1-800-609-4884 (toll free) OR mail requests to: PharmaCare, Box 9652 Stn Prov Govt, Victoria, BC V8W 9P4 This facsimile is doctor-patient privileged and contains confidential information intended only for PharmaCare. Any other distribution, copying or disclosure is strictly prohibited.

MISDIRECTED across the front of the form and fax toll-free to 1-800-609-4884, then destroy the pages received in error.

If you have received this fax in error, please write

If PharmaCare approves this Special Authority request, approval is granted solely for the purpose of covering prescription costs. PharmaCare approval does not indicate that the requested device is, or is not, suitable for any specific patient or condition.

Forms with information missing will be returned for completion. If no prescriber fax or mailing address is provided, PharmaCare will be unable to return a response.

SECTION 1 – PRESCRIBER'S INFORMATION	SECTION 2 - PATIENT INFO	ORMATION	
Prescriber's Name and Mailing Address	Patient (Family) Name	Patient (Family) Name	
	Patient (Given) Name(s)		
College ID (use ONLY College ID number) Phone Number (include area cod	e) Date of Birth (yyyy / mm / dd)	Date of Application (yyyy / mm / dd)	
CRITICAL FOR A TIMELY RESPONSE Prescriber's Fax Number	CRITICAL FOR PROCESSING	sonal Health Number (PHN)	
SECTION 3 - MEDICATION REQUESTED	SACUBITR	il + valsartan: 9901-0289	
sacubitril-valsartan (24 mg/26 mg, 49 mg/51 mg and 97 m	g/103 mg tablets)		
SECTION 4 – CRITERIA FOR INDEFINITE COVERAGE			
Approval subject to ALL of the criteria below being met (mark boxes and complete blanks as applicable):			
Prescriber authorizing this request is an internal medicine specialist or cardiologist.			
Patient has heart failure with New York Heart Association Class II to stable dose:	Il symptoms that have persisted despite at least	t four weeks of treatment at the optimum	
$\hfill \Box$ Of a beta-blocker and other recommended therapies, including	ng an aldosterone antagonist (if tolerable).		
Of an angiotensin-converting enzyme inhibitor (ACEI) or an a	ngiotensin II receptor antagonist (ARB).		
Name of ACEI or ARB trialed:			
Duration of trial of prior ACEI or ARB (minimum 4 weeks req	uired):		
☐ Provide left ventricular ejection fraction (LVEF) (≤40%):			
Sacubitril-valsartan should be administered in place of an ACEI or an ARB.			
Report all adverse events to the post-market surveillance prog		5-234-2345 (health professionals only).	
Personal information on this form is collected under the authority of, and in accordance with, the <i>British Columbia Pharmaceutical Services Act</i> 22(1) and <i>Freedom of Information and Protection of Privacy Act</i> 26 (a),(c),(e). The information is being collected for the purposes of (a) administering the PharmaCare program, (b) analyzing, planning and evaluating the Special Authority and other Ministry programs and (c) to manage and plan for the health system generally. If you have any questions about the collection of this information, call Health Insurance BC from Vancouver at 1-604-683-7151 or from elsewhere in BC toll free at		t that the purpose of releasing their obtain Special Authority for prescription	
1-800-663-7100 and ask to consult a pharmacist concerning the Special Authority pro	cess. Eligible Specialist's Signature (Mandatory)	
PharmaCare may request additional documentation to support this Special Aut Actual reimbursement is subject to the rules of a patient's PharmaCare plan, inc		any other applicable PharmaCare pricing policy.	
PHARMACARE USE ONLY			
STATUS	EFFECTIVE DATE (YYYY / MM / DD)	DURATION OF APPROVAL	