

PHARMACARE SPECIAL AUTHORITY REQUEST SATRALIZUMAB FOR NEUROMYELITIS OPTICA **SPECTRUM DISORDER (NMOSD)**

HLTH 5812 Rev. 2023/10/31

○ INITIAL Complete sections 1, 2, & 3		RENEWAL Complete sections 1, 2, & 4				
	Complete se	ctions 1, 2, & 3		Complete so	ections	1, 2, & 4
For up-to-date criteria and for Fax requests to 1-800-609-4884 This facsimile is doctor-patient priv copying or disclosure is strictly prol	(toll free) OR mail requests to: Plilieged and contains confidential info	harmaCare, Box 9652	Stn Prov Govt, Vic	ctoria, BC V8W 9P4	MISD toll-fr	have received this fax in error, please write IRECTED across the front of the form and fax ee to 1-800-609-4884, then destroy the pages yed in error.
If PharmaCare approves this Specia	l Authority request, approval is grar cate that the requested device is, or					
• • • • • • • • • • • • • • • • • • • •	•	•			PharmaC	are will be unable to return a response.
		·	_	-		•
SECTION 1 - NEUROLO		ı	SECTION 2 – PATIENT INFORMATION			
Neurologist's Name and Maili	ng Address		Patient (Fai	mily) Name		
			Patient (Giv	ven) Name(s)		
College ID (use ONLY College ID number) Phone Number (include area c		clude area code)	Date of Birt	Date of Birth (yyyy / mm / dd) Date of Applica		Date of Application (yyyy / mm / dd)
	Neurologist's Fax Number				Personal	l Health Number (PHN)
CRITICAL FOR A TIMELY RESPONSE			PROCESS			
SECTION 3 - INITIAL CO	OVERAGE CRITERIA FO	R SATRALIZUM	AB: 1 YEAR			SATRALIZUMAB: 9901-0440
Dosage: 120mg SC at 0, 2 a	nd 4 weeks, followed by 120n	ng SC every 4 week	s for maintenan	ce		
	aporin 4 (AQP4) seropositive NA					
	ist with expertise in the diagno					
		_		the 3-month period	l immedia	tely preceding satralizumab initiation.
	, da	te		·		
PLUS one of the following: A. patient has had tr	eatment failure resulting in at l	east one relapse of N	MOSD within the	e previous 12 month	ns despite	a trial of optimally dosed
rituximab OR toci	izumab					
	erate or has contraindications t	o both rituximab ANI	D tocilizumab, lea	ading to discontinua	ation or in	ability to use rituximab and tocilizumab.
NAME OF PREVIOUSLY TRIED THERAPIE	S DOSE	DURATION OF TRIAL (MONTHS)		DETAILS (FAILURE, CONTRAINDIC	OF OUTCOM	
THE VIOUS ET THE STITLING TO	5 5052	THINE (MONTHS)	○ Failure	O Intolerance		ntraindication
1 Rituximab			Specify:			
			○ Failure	OIntolerance	○ Co	ntraindication
2 Tocilizumab			Specify:			

Please complete additional information on page 2 >>

PHARMACARE USE ONLY					
TATUS	EFFECTIVE DATE	DURATION OF THERAPY / TERMINATION DATE			

SATRALIZUMAB FOR NEUROMYELITIS OPTICA SPECTRUM DISORDER (NMOSD)

PATIENT NAME		PHN	DATE (YYYY / MM / DD)
SECTION 4 – RENEWAL COVER	AGE CRITERIA FOR SATRALIZU	JMAB: 1 YEAR	
Dosage: 120mg SC every 4 weeks			
Prescribed by a neurologist with expe	ertise in the diagnosis and management o	of NMOSD.	
Patient has maintained an EDSS score	e of less than 8 points taken within the 3-1	month period imme	ediately preceding the renewal request.
Most recent EDSS score	, date	·	
ECTION E ADDITIONAL NOT	EC		
ECTION 5 - ADDITIONAL NOT	E2		
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		information to	sed with the patient that the purpose of releasing their to PharmaCare is to obtain Special Authority for prescription the purposes set out here.
system generally. If you have any questions about Health Insurance BC from Vancouver at 1-604-6	out the collection of this information, call		
1-800-663-7100 and ask to consult a pharmacist concerning the Special Authority process.		Navvala siat/a Ciara	. (4)

 $Pharma Care\ may\ request\ additional\ documentation\ to\ support\ this\ Special\ Authority\ request.$

Actual reimbursement is subject to the rules of a patient's PharmaCare plan, including any annual deductible requirement, and to any other applicable PharmaCare pricing policy.