

## CHRONIC MEDICINE BENEFIT APPLICATION FORM – 2011

*(Applies to Nedgroup Traditional Plus members only)*

- Please complete the application in black ink
- One application form must be completed per patient
- Please attach a copy of the Dr's prescription to the application form (original not required)
- Applications will not be processed unless the appropriate sections are completed and relevant documents are attached. The completed and signed application form may be faxed to **086 679 1579**, emailed to [onecare@scriptpharm.co.za](mailto:onecare@scriptpharm.co.za) or posted to Scriptpharm Risk Management, P.O. Box 653590, Benmore, 2010
- Clinical entry criteria must be met before medication for Prescribed Minimum Benefit (PMB) or chronic conditions will be authorised. See Section H.
- **Please note that Chronic medication approved but not claimed for at least 6 consecutive months will be terminated and the member will have to re-apply for the benefit with all the relevant tests and a new application form.**
- **Contact the call centre on 010 591 0150 for further assistance**

### SECTION A. PRINCIPAL MEMBER'S DETAILS

Membership Number  Scheme and Option

Surname

Title  Initials  Date of Birth

Telephone numbers Home  (  ) Work  (  )

Cell  Fax  (  )

Postal Address   
  
 Postal code

Email Address (will be treated as private)

### SECTION B. PATIENT'S DETAILS

Surname  Title

Full first name  Dependant Code

Date of Birth             Gender (M/F)

Telephone numbers Home  (  ) Work  (  )

Cell  Fax  (  )

Email Address (will be treated as private)

Please **circle** the preferred method of communication (if patient is under the age of 16 years, communication will be sent to the main member)

Email

Fax

Post

**Please ensure that relevant details have been provided for the communication option selected**

### SECTION C. DECLARATION BY PATIENT (or member if patient is a minor)

I hereby authorise my doctor to furnish and/or disclose any relevant clinical information required to review my application. I understand that the application is subject to formulary guidelines as well as Scheme rules. I also understand that generic equivalents will be authorised where applicable and co-payments will apply if I choose not to accept the substitution.

Patient signature (unless a minor)

Date

Patient name and surname

Membership number

**SECTION D. CARDIOVASCULAR RISK (to be completed by doctor when applying for PMB benefits for hypertension, hyperlipidaemia, diabetes mellitus type 2 and cardiac failure)**

Weight in kg  Patient height in metres  Body Mass Index

Does the patient smoke? Yes/No

Is microalbuminuria present or is the GFR less than 60ml/min? Yes/No

If there is target organ damage and/or cardiovascular disease, please tick the appropriate box

- |  |  |   |
|--|--|---|
| <input type="checkbox"/> Angina        | <input type="checkbox"/> Myocardial Infarction | <input type="checkbox"/> Hypertensive Retinopathy     |
| <input type="checkbox"/> Heart Failure | <input type="checkbox"/> Prior Stenting        | <input type="checkbox"/> Left Ventricular Hypertrophy |
| <input type="checkbox"/> Prior CABG    | <input type="checkbox"/> Cardiomyopathy        | <input type="checkbox"/> Peripheral Arterial Disease  |
| <input type="checkbox"/> Stroke        | <input type="checkbox"/> Chronic Renal Disease | <input type="checkbox"/> Transient Ischaemic Attack   |

For cardiac failure, please provide either the NYHA classification: Class \_\_\_\_\_, or the stage of cardiac failure according to the American College of Cardiology/ American Heart Association Task Force on Practice Guidelines: Stage \_\_\_\_\_

**SECTION E. APPLICATION FOR HYPERTENSION**  
*Please complete in conjunction with Section D*

**A specialist must complete this section for patients below the age of 30 years diagnosed with hypertension**

1. Current blood pressure \_\_\_\_\_ / \_\_\_\_\_ mmHg

2. When did the patient commence drug therapy for hypertension? 

Y	Y	Y	Y	M	M	D	D
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3. For all newly diagnosed patients and those diagnosed in the last 6 months, please supply the 2 initial blood pressure readings (before drug therapy), performed at least 2 weeks apart

Date  /  mmHg      Date  /  mmHg

4. Please provide additional clinical information if there are compelling indications for use of drug classes that are not first or second line therapy, such as Angiotensin Receptor Blockers.

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**SECTION F. APPLICATION FOR HYPERLIPIDAEMIA**  
*Please complete in conjunction with Section D*

**Please attach a copy of a recent full lipogram.**

1. Please list the signs of Familial Hyperlipidaemia, if present \_\_\_\_\_

2. Is there a family history of premature arteriosclerotic disease? Yes/ No   
 If the answer is YES, please provide the following details:

	Father	Mother	Sibling
Description of event			
Age at time of first event			

3. When did your patient commence drug therapy for hyperlipidaemia? 

Y	Y	Y	Y	M	M	D	D
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4. In terms of the European Guidelines adopted by the South African Heart Association, patients falling in the following categories are not required to be risk scored.  
 Please provide supporting clinical evidence or pathology results to confirm the health status of the patient.

1. Established atherosclerosis:
  - a. Coronary Heart Disease
  - b. Cerebrovascular atherosclerotic disease
  - c. Peripheral vascular disease
2. Diabetes Type 2
3. Diabetes Type 1 with microalbuminuria or proteinuria

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5. For patients with **primary hyperlipidaemia**, please assess your patient's risk using the following table. Kindly indicate the score by marking the appropriate percentage risk

Estimate of 10-year risk for **MEN**

Age (years)	Points
20-34	-9
35-39	-4
40-44	0
45-49	3
50-54	6
55-59	8
60-64	10
65-69	11
70-74	12
75-79	13

Estimate of 10-year risk for **WOMEN**

Age (years)	Points
20-34	-7
35-39	-3
40-44	0
45-49	3
50-54	6
55-59	8
60-64	10
65-69	12
70-74	14
75-79	16

Total Cholesterol (mmol/ L)	Points Age (years)				
	20-39	40-49	50-59	60-69	70-79
<4	0	0	0	0	0
4.1-5	4	3	2	1	0
5.1-6.2	7	5	3	1	0
6.21-7.2	9	6	4	2	1
≥7.2	11	8	5	3	1

Total Cholesterol (mmol/ L)	Points Age (years)				
	20-39	40-49	50-59	60-69	70-79
<4	0	0	0	0	0
4.1-5	4	3	2	1	1
5.1-6.2	8	6	4	2	1
6.21-7.2	11	8	5	3	2
≥7.2	13	10	7	4	2

	Points Age (years)				
	20-39	40-49	50-59	60-69	70-79
Non-smoker	0	0	0	0	0
Smoker	8	5	3	1	1

	Points Age (years)				
	20-39	40-49	50-59	60-69	70-79
Non-smoker	0	0	0	0	0
Smoker	9	7	4	2	1

Estimate of 10-year risk for <b>MEN</b>	
HDL (mmol/L)	Points
≥1.6	-1
1.30-1.59	0
1.00-1.29	1
<1	2

Estimate of 10-year risk for <b>WOMEN</b>	
HDL (mmol/L)	Points
≥1.6	-1
1.30-1.59	0
1.00-1.29	1
<1	2

Systolic BP (mmHg)	Points	
	If untreated	If treated
<120	0	0
120-129	0	1
130-139	1	2
140-159	1	2
≥160	2	3

Systolic BP (mmHg)	Points	
	If untreated	If treated
<120	0	0
120-129	1	3
130-139	2	4
140-159	3	5
≥160	4	6

Estimate of 10-year risk for <b>MEN</b>		
	Total Points	10-year risk %
	<0	<1
	0	1
	1	1
	2	1
	3	1
	4	1
	5	2
	6	2
	7	3
	8	4
	9	5
	10	6
	11	8
	12	10
	13	12
	14	16
	15	20
	16	25
	≥ 17	≥ 30

10-year risk \_\_\_\_%

Estimate of 10-year risk for <b>WOMEN</b>		
	Total Points	10-year risk %
	<9	<1
	9	1
	10	1
	11	1
	12	1
	13	2
	14	2
	15	3
	16	4
	17	5
	18	6
	19	8
	20	11
	21	14
	22	17
	23	22
	24	27
	≥ 25	≥ 30

10-year risk \_\_\_\_%

**Framingham scoring system for calculating the 10-year risk of major coronary events in adults without diabetes.**

HDL denotes high-density lipoprotein cholesterol & BP blood pressure. All age ranges are given in years. Reprinted from National Institutes of Health, National Heart, Lung and Blood Institute. Third Report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation and Treatment of High blood cholesterol in Adults (Adult Treatment Panel III). Executive Summary. NIH Publication No. 01-3670; May 2001.

**6. Based on the information supplied in Section F:**

- For patients **below** the age of **60** years: Does your patient have a 20% or greater risk of a coronary event in the next ten years? (Please circle Yes/No)      **Yes**                      **No**
- For patients **above** the age of **60**: Does your patient have a 30% or greater risk of a coronary event in the next ten years? (Please circle Yes/No)      **Yes**                      **No**

We acknowledge that there are limitations to the Framingham Risk Assessment Score Chart. In order to assist with a funding decision, please motivate if you feel that your patient is negatively impacted by these limitations.

The PMB benefit will *not* provide cover in patients with less than a 20% (<60 years) or 30%(>60 years) risk of a coronary artery event within the next ten years. This is based on the local and international treatment guidelines and is in line with the Medical Scheme Council Clinical Algorithm. This is a funding decision, to ensure the long term sustainability of this benefit and does not in any way question your clinical decision.

**SECTION G: APPLICATION FOR OSTEOPOROSIS (to be completed by Medical Practitioner. Please attach a BMD report)**

Osteoporotic fracture: (Please circle Yes or No) <b>Yes</b> <b>No</b>	If yes, please supply date of most recent fracture:	Y	Y	Y	Y	M	M	D	D
Please indicate fracture location/s:									



**SECTION H. PRESCRIBED MINIMUM BENEFITS: CLINICAL ENTRY CRITERIA**

1. Please note that your application will not be processed if the requested information is not supplied
  2. Some conditions may require completion of the form by a relevant specialist
  3. Each time you register for **a new chronic disease**, the information in the following table is required.
- Once registered for a chronic condition, you may be required to submit further documentation if your medication is changed.

	<b>SUBMISSION REQUIREMENTS</b>
a. Application for a change in medicine where you are currently registered for the same condition.	Section A, B and C and a copy of a valid prescription.
b. Application for medication for a second condition where you have already registered for a first.	Complete application form including clinical criteria and copy of valid prescription.
c. If the condition applied for was approved by your previous medical scheme, a report from your doctor stating the name of the condition, medication and duration of treatment is required.	Section A, B and C, a letter of motivation from the prescriber and a copy of a valid prescription.

<b>PMB CONDITION</b>	<b>CLINICAL ENTRY CRITERIA</b>
<b>Addison's Disease</b>	<ol style="list-style-type: none"> <li>1. Serum cortisol levels</li> <li>2. ACTH stimulation test to distinguish primary from secondary adrenal insufficiency. The PMB is only applicable to primary Addison's disease</li> <li>3. A specialist physician, paediatrician or endocrinologist must make the diagnosis.</li> </ol>
<b>Asthma</b>	<ol style="list-style-type: none"> <li>1. A lung function test for adults and children older than 7 years</li> <li>2. The South African Treatment Guidelines for asthma will be used to assess all applications</li> <li>3. Applications for leukotriene inhibitors (e.g. Singulair<sup>®</sup>) must be supported by a pre- and post lung function test to substantiate the additional benefit and must be from a Pulmonologist.</li> </ol>
<b>Bipolar Mood Disorder</b>	A psychiatrist prescription and written diagnosis are required.
<b>Bronchiectasis</b>	Please attach a report based on the findings of a radiological examination (Chest X-ray or CT scan)
<b>Cardiac Failure</b>	<ol style="list-style-type: none"> <li>1. Please indicate the level of functional incapacity according to the New York Heart Association's classification and/or</li> <li>2. The stage of cardiac failure according to the American College of Cardiology/ American Heart Association Task Force on Practice Guidelines (February 2002)</li> </ol> Please record level/stage in Section D
<b>Cardiomyopathy</b>	The diagnosis must be confirmed by a specialist physician or cardiologist
<b>Chronic Obstructive Pulmonary Disease (COPD)</b>	Please attach a lung function test. The REF (risk equalisation fund) criteria are in line with the GOLD classification
<b>Chronic Renal Disease</b>	<ol style="list-style-type: none"> <li>1. A specialist physician must complete the application</li> <li>2. Indicate the creatinine clearance</li> <li>3. When applying for erythropoetin, a report indicating haemoglobin, T<sub>sat</sub> and ferritin levels must be provided. Please also state whether the patient is currently on or off drug therapy</li> <li>4. A report indicating T<sub>sat</sub> and ferritin must be provided when applying for iron supplementation</li> </ol>
<b>Coronary Artery Disease</b>	Please attach a copy of the stress or exercise ECG report confirming the diagnosis of coronary artery disease
<b>Crohn's Disease</b>	The application form must be completed by a gastroenterologist or specialist physician. If the condition is managed by a general practitioner, a gastroenterologist must confirm the diagnosis

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<b>PMB CONDITION</b>	<b>CLINICAL ENTRY CRITERIA</b>
<b>Diabetes Insipidus</b>	<ol style="list-style-type: none"> <li>1. An endocrinologist, specialist physician, paediatrician, neurologist or neurosurgeon must complete the application form</li> <li>2. The results of a water deprivation test are required</li> </ol>
<b>Diabetes Mellitus Type I</b>	Application form must be completed by a medical practitioner
<b>Diabetes Mellitus Type II</b>	Section D must be completed by a medical practitioner. Blood tests required.
<b>Dysrhythmias</b>	The medical practitioner must indicate the ICD 10 code. The PMB chronic benefit only provides cover for chronic atrial fibrillation and flutter (I48) and ventricular tachycardia (I47.2)
<b>Epilepsy</b>	<ol style="list-style-type: none"> <li>1. Please attach a detailed seizure history</li> <li>2. Please attach an EEG report confirming the diagnosis of epilepsy</li> </ol>
<b>Glaucoma (open and closed angle)</b>	Please provide the intra-ocular pressure at diagnosis. This is only required for newly diagnosed patients
<b>Haemophilia</b>	Haemophilia A: Please provide the Factor VIII level as a % of normal Haemophilia B: Please provide the Factor IX level as a % of normal
<b>Hyperlipidaemia</b>	Please attach a copy of the diagnosing (for primary hyperlipidaemia) or current (for secondary hyperlipidaemia) lipogram. The medical practitioner must complete Sections D and F of the application form.
<b>Hypertension</b>	Section D and E of the application form must be completed by the medical practitioner
<b>Hypothyroidism</b>	Please attach the diagnostic report that confirms the initial diagnosis of hypothyroidism
<b>Multiple Sclerosis</b>	<ol style="list-style-type: none"> <li>1. A specialist physician or neurologist must complete the application form and indicate the specific type of multiple sclerosis</li> <li>2. Please provide the following information when applying for chronic medicine benefits for inteferon: <ol style="list-style-type: none"> <li>a) Extended disability status score (EDSS)</li> <li>b) Relapsing-remitting history</li> <li>c) Number of relapses requiring IV cortisone treatment</li> </ol> </li> </ol>
<b>Parkinson's Disease</b>	Applications for non-formulary products will only be considered if prescribed by a neurologist, or if the application is supported by a neurologist's motivation
<b>Rheumatoid Arthritis</b>	<ol style="list-style-type: none"> <li>1. Copies of the relevant blood test reports and supportive clinical history confirming the diagnosis of rheumatoid arthritis are required</li> <li>2. Applications for COXIBs must be supported by a motivation indicating the risk factors considered for their use over conventional anti-inflammatories</li> <li>3. Applications for anti-inflammatories as monotherapy <b>MUST</b> be motivated by a rheumatologist</li> </ol>
<b>Schizophrenia</b>	A psychiatrist prescription and written diagnosis is required
<b>Systemic Lupus Erythematosus (SLE)</b>	A rheumatologist, specialist physician or paediatrician must complete the application form and indicate the diagnostic criteria used
<b>Ulcerative Colitis</b>	A gastroenterologist or specialist physician must complete the application form. If the condition is managed by a general practitioner, a gastroenterologist or specialist physician must confirm the diagnosis

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<b>SECTION I. NON-PRESCRIBED MINIMUM BENEFITS CHRONIC DISEASES</b> <i>(subject to a medicines formulary)</i>	
<b>CHRONIC CONDITION</b>	<b>CLINICAL ENTRY CRITERIA REQUIREMENTS</b>
<b>Acne (Cystic nodular only)</b>	For isotretinoin therapy, the patient's weight, date of commencement with treatment and duration of therapy is required. A dermatologist must initiate therapy
<b>Allergic Rhinitis</b>	Only covered in children under the age of 12 years, or in patients on concurrent asthma therapy. A motivation or a specialist prescription is required for the combined use of inhaled nasal corticosteroids and antihistamines.
<b>Alzheimer's Type Dementia</b>	Please submit the results of a mini-mental state examination (MMSE)
<b>Anxiety</b>	Only reviewed if member is approved for a PMB psychiatric condition
<b>Attention Deficit Disorder (ADHD or ADD)</b>	A paediatrician, psychiatrist or neurologist must complete the application form. This condition will only be covered in patients under the age of 18 years.
<b>Deep vein thrombosis</b>	No clinical entry criteria
<b>Major Depression</b>	Generic fluoxetine or citalopram will be funded as first-line therapy from a GP for 6 months, pending review from a psychiatrist. An initial psychiatrist's prescription is required for all other anti-depressants and mood stabilisers.
<b>Gastro-oesophageal Reflux Disease (GORD)</b>	GORD will only be funded if the patient suffers from a GIT-related PMB condition, or is on oral corticosteroid treatment for another PMB condition. A gastroscopy report, including the Los Angeles Grading is required. Generic omeprazole, cimetidine or ranitidine will be funded. Please submit a detailed, clinically relevant motivation for other products.
<b>Gout</b>	Only patients whose Gout is associated with Diabetes and/or Hypertension
<b>Hyperthyroidism</b>	Submit T3, T4 and TSH levels. Funding will be authorised for 6 months pending review
<b>Hyperpituitarism</b>	A specialist physician or endocrinologist must complete the application form. Basal/stimulation test results required
<b>Hypopituitarism</b>	A specialist physician or endocrinologist must complete the application form. Basal/stimulation test results required
<b>Hypoparathyroidism</b>	A specialist physician or endocrinologist must complete the application form. Calcium, Phosphate and parathyroid hormone levels required.
<b>Hypotension</b>	A pre and post medication blood pressure reading must be submitted
<b>Insomnia</b>	Only if linked to another PMB psych condition (Bipolar or Schizophrenia)
<b>Menopause (HRT)</b>	Tibolone will only be authorised on motivation from a gynaecologist. Motivation required if patients less than 40 years old
<b>Migraine</b>	Only prophylaxis will be covered
<b>Osteoarthritis</b>	Applications for COXIBs must be supported by a motivation indicating the risk factors considered for their use over conventional anti-inflammatories
<b>Osteoporosis</b>	<ol style="list-style-type: none"> <li>1. Applications must include a DEXA bone mineral density scan (BMD) report</li> <li>2. A short report on additional risk factors must be included (e.g. previous fractures, family history, long term oral corticosteroid use). Please complete Section G</li> <li>3. An endocrinologist motivation is required for males, females under the age of 30, and children.</li> </ol>
<b>Psoriasis</b>	No clinical entry criteria

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