

C. Diagnosis

3. What is the primary diagnosis in relation to the requested equipment?

- | | | |
|--|--|--|
| <input type="checkbox"/> Bronchiectasis | <input type="checkbox"/> COPD | <input type="checkbox"/> Interstitial lung disease |
| <input type="checkbox"/> Cardiac failure intractable | <input type="checkbox"/> COVID-19-pneumonitis/long COVID | <input type="checkbox"/> Pulmonary fibrosis |
| <input type="checkbox"/> Congenital cardiac disease | <input type="checkbox"/> Cystic fibrosis | <input type="checkbox"/> Pulmonary hypertension |
| <input type="checkbox"/> Other <input type="text"/> | | |

4. Provide other relevant diagnosis/co-morbidities

D. Equipment specification

5. Select one stationary oxygen concentrator and specify flow rate: Select ONE option

- Oxygen concentrator standard flow (0-5 L/min). Flow rate (L/min)
- Oxygen concentrator high flow (6-10 L/min). Flow rate (L/min)
- No oxygen concentrator requested

6. Select C size cylinders and specify flow rate: Select ONE option

- Portable oxygen C cylinders x2 with standard regulator x1. Flow rate (L/min)
- Portable oxygen C cylinders x2 with conserver regulator x1. Flow rate (L/min)
- No portable cylinders requested

7. Does the person use continuous oxygen (≥ 16 hours/day) AND reside > 2 hours from their closest hospital?

- No
- Yes – provide D Cylinder. Flow rate (L/min)

E. Current supplier of home oxygen

8. Does the person currently receive supply of home oxygen (e.g. hospital discharge supply)? Select ONE option

- No
- Yes-Initial oxygen supplied by BOC
- Yes-Initial oxygen supplied by Supagas

F. Short term oxygen therapy discharge information

9. Confirm ALL of the following have been addressed:

- The person is in hospital with an acute illness and in the recovery phase, nearing their date of discharge.
Provide estimated or known discharge date from acute facility
- The first month's supply of oxygen has been funded by the discharging service

G. Stability, compliance and ongoing follow-up

10. Confirm ALL of the following have been addressed:

- The person is aware that they will not be eligible for funding if they smoke
- Recommended oxygen equipment is compatible with the person's living environment
- The person is aware that data regarding oxygen therapy usage will be collected by the supplier and can be obtained by the prescriber and EnableNSW
- Review for long term oxygen has been arranged. Provide follow-up date

H. Eligibility: oxygen concentrator

11. Select ONE of the oxygen usage requirements and complete the relevant criteria below:

Prescription is ≥ 16 hours/day

Choose one criteria, ensure stable ABG is attached and provide PaO₂

Daytime PaO₂ ≤ 55 mmHg **OR**

Daytime PaO₂ 56–59 mmHg plus written evidence of significant end-organ damage due to one or more of the following: pulmonary hypertension, right heart failure, polycythaemia or other.

Specify evidence below:

OR

Prescription is ≥ 6 hours/day (for nocturnal hypoxaemia)

Confirm one from EACH category (Dx-Diagnostic criteria, Evidence of improvement and Positive Airway Pressure (PAP) intolerance). Attach any supporting test results, ABG, and clinical reports/letters

Dx: SpO₂ $\leq 88\%$ for $\geq 30\%$ (attach technical and physician report of polysomnogram or nocturnal oximetry)

OR

Dx: SpO₂ $\leq 80\%$ for $\geq 10\%$ of sleep time (attach technical and physician report of polysomnogram or nocturnal oximetry)

AND

Evidence of improvement: provide technical and physician report of polysomnogram or nocturnal/continuous oximetry demonstrating objective improvement in SpO₂ (attached)

AND

PAP intolerance: Is the person unable to tolerate PAP and prescribed oxygen as an alternative treatment for sleep-disordered breathing?

N/A

Yes-provide letter of justification

I. Improvement and stability on titrated oxygen flow rate

12. Confirm stability and safety on prescribed oxygen:

The person's oxygen flow rate has been adequately titrated to ensure SpO₂ is maintained within a safe target range for the person, and to avoid worsening hypercapnia

J. Eligibility: portable cylinder oxygen (C cylinders)

13. Confirm ONE of the following and complete any relevant sections below:

Prescription is ≥ 16 hours/day

Prescription is <16 hours/day

If <16 hours/day selected, confirm ALL the following and attach clinical letter and 6 minute walk tests

Diagnosis of interstitial lung disease or other non-COPD lung disease

OR

Respiratory diagnosis with evidence of hypoxia-related sequelae (polycythaemia, right heart failure, pulmonary hypertension), provided in a clinical letter of justification.

AND

Evidence of significant desaturation during exercise (SpO₂ $<88\%$) while breathing room air

AND

Distance walked in 6 minute walk test while on oxygen improves by $\geq 25m$ OR by $> 50\%$ in people with baseline 6 minute walk distance $< 50m$

N/A -Portable Cylinder Oxygen is not requested

K. Funding portable oxygen cylinder refills

14. Confirm the following for portable oxygen cylinder requests:

- The person is aware of and willing to partially fund the therapy including charges for portable C cylinder refills and delivery charges
- N/A - Portable Cylinder Oxygen is not requested

L. Community safety, training and emergency plan

15. Confirm ALL of the following three criteria demonstrating adequate community safety, carer training and provision of an emergency plan have been addressed:

- A risk assessment has been conducted and documented, and the person can be safely managed on the prescribed equipment in the community
- The person and family/carer/s have received adequate training, and have acknowledged the risks and responsibility for safely managing the person and the equipment in the community
- An individual care plan and an emergency plan have been documented and communicated to the person and their family/carer/s, to manage clinical and equipment emergencies and to allow the person to live safely in the community

M. Ongoing monitoring and assessment

16. Provide the details of the eligible clinician/prescriber who will continue to monitor the person: *Select ONE option*

- The prescriber for this request (Respiratory/Palliative Care Physician) will assess and monitor the person's condition
- A different eligible prescriber (Respiratory/Palliative Care Physician) will assess and monitor the person's condition

Provide name, qualification, phone number, email address and clinical service:

- An eligible Nurse Practitioner will assess and monitor the person's condition, working in collaboration with a Respiratory or Palliative Care Physician

Provide name, qualification, phone number, email address and clinical service, and name of the Respiratory or Palliative Care Physician:

- Requests from other prescribers, such as general practitioners or physicians, will only be considered in rural or remote areas, where an eligible prescriber (Respiratory Physician, Palliative Care Physician or Respiratory Nurse Practitioner) is unavailable within the health service/ Local Health District.

If this is the case, with each application the prescriber must provide a letter:

- Outlining the reasons why an eligible prescriber is not available **AND**

Provide name, qualification, phone number, email address of the clinician responsible for follow-up and ongoing respiratory care of the person:

Go to next page and complete Section N. Prescriber eligibility and declaration

N. Prescriber eligibility and declaration

17. Prescriber eligibility

Confirm you have assessed the person and have the qualification and level of experience to prescribe this equipment in line with the relevant [EnableNSW Funding Criteria](#) and [Professional Criteria for Prescribers](#).

Yes

18. Prescriber declaration

I confirm the following:

- The person/carer agrees with this request
- A copy of this request will be provided to the person/carer
- As a health professional, I cannot also be the equipment supplier for the same request. This may include but is not limited to a personal or professional relationship with or material interest in the supplier or manufacturer of the equipment listed on this request

I declare that:

- I have the qualification and experience to prescribe this equipment or, I have been supervised by an eligible EnableNSW prescriber for this type of equipment
- All information I have supplied on this application is true and correct to the best of my knowledge at the time of assessment

Prescriber information:

Prescriber name	<input type="text"/>			
Place of work	<input type="text"/>			
Address	<input type="text"/>		State <input type="text"/>	Postcode <input type="text"/>
Qualification	<input type="text"/>	AHPRA registration number	<input type="text"/>	
Phone number	(<input type="text"/>) <input type="text"/>	Email	<input type="text"/>	
Signature	<input type="text"/>		Date	<input type="text" value="D D/M M/YYYY"/>

19. Other contacts (optional)

Complete this question if you would like to provide details of any other relevant health professionals who will be involved with the management and monitoring of the person's condition

Other contact 1

Name	<input type="text"/>			
Place of work	<input type="text"/>			
Address	<input type="text"/>		State <input type="text"/>	Postcode <input type="text"/>
Qualification	<input type="text"/>	AHPRA registration number	<input type="text"/>	
Phone number	(<input type="text"/>) <input type="text"/>	Email	<input type="text"/>	

Other contact 2

Name	<input type="text"/>			
Place of work	<input type="text"/>			
Address	<input type="text"/>		State <input type="text"/>	Postcode <input type="text"/>
Qualification	<input type="text"/>	AHPRA registration number	<input type="text"/>	
Phone number	(<input type="text"/>) <input type="text"/>	Email	<input type="text"/>	

Submitting this request

Submit this form and any relevant clinical documentation to enable@health.nsw.gov.au, please include the following in your subject line **Equipment type_Person name_Date submitted** i.e *Oxygen_Adult_STOT_request_John Smith_01.01.2022*