

1
2
3 **British Thoracic Society Guideline for oxygen use in adults**
4 **in healthcare and emergency settings**

5
6 **Concise Guideline: Public consultation copy**
7 **30/11/2015**

8
9 *B R O'Driscoll, L Howard, J Earis, V Mak,*
10 *on behalf of the BTS Emergency Oxygen Guideline Group*
11

12
13
14
15
16
17 **Available for public consultation from 30 November 2015 to 18 January 2016.**
18

19
20
21 Contact: British Thoracic Society, 17 Doughty St, London WC1N 2PL

22 sally.welham@brit-thoracic.org.uk

23 louise.preston@brit-thoracic.org.uk
24
25
26

27 *The concise version of the BTS Guideline should be read in conjunction with the full version –*
28 *also available for public consultation at:*

29 [https://www.brit-thoracic.org.uk/guidelines-and-quality-standards/emergency-oxygen-use-](https://www.brit-thoracic.org.uk/guidelines-and-quality-standards/emergency-oxygen-use-in-adult-patients-guideline/)
30 [in-adult-patients-guideline/](https://www.brit-thoracic.org.uk/guidelines-and-quality-standards/emergency-oxygen-use-in-adult-patients-guideline/)
31

32

33 *This guideline replaces the 2008 British Thoracic Society guideline for emergency oxygen use.*
34 O'Driscoll BR, Howard LS, Davison AG; British Thoracic Society. BTS guideline for emergency
35 oxygen use in adult patients. Thorax. 2008 Oct;63 Suppl 6:vi1-68.[1]
36
37

38

39

40 *BTS Emergency Oxygen Guideline Group 2015*

41

42 Dr Sabrina Bajwah

43 Professor Richard Beasley

44 Dr Katrina Curtis

45 Professor Anthony Davison

46 Dr Alistair Dorward

47 Dr Chris Dyer

48 Ms Angela Evans

49 Ms Lucy Falconer

50 Ms Clare Fitzpatrick

51 Dr Simon Gibbs

52 Dr Kim Hinshaw

53 Dr Robin Howard

54 Dr Binita Kane

55 Dr Jeff Keep

56 Ms Carol Kelly

57 Mr Hasanin Khachi

58 Dr Muhammed Asad Iqbal Khan

59 Dr Roop Kishen

60 Ms Leigh Mansfield

61 Dr Bruce Martin

62 Dr Fionna Moore

63 Dr Duncan Powrie

64 Dr Louise Restrick

65 Professor Christine Roffe

66 Dr Mervyn Singer

67 Dr Jasmeet Soar

68 Dr Iain Small

69 Ms Lisa Ward

70 Mr David Whitmore

71 Professor Wisia Wedzicha

72 Dr Meme Wijesinghe

73

74

75

76

77	Contents
78	
79	
80	Executive Summary of the Guideline
81	
82	Key changes since the 2008 Guideline was published
83	
84	1. Introduction
85	1.1 Aim of the Guideline
86	1.2 Intended users of the guideline and target patient populations
87	1.3 Areas covered by the guideline
88	1.4 Areas not covered by the guideline
89	1.5 Limitations of the guideline
90	
91	2. Methodology of Guideline production
92	2.1 Establishment of guideline team
93	2.2 Planned review and updating of the guideline
94	2.3 Declarations of interest
95	
96	
97	3. Summary of recommendations and good practice points:
98	A. Achieving desirable oxygen saturation ranges in acute illness
99	B. Clinical and laboratory assessment of hypoxaemia and hypercapnia
100	C. Arterial and arterialised blood gases
101	D. Initial oxygen therapy; initial choice of equipment
102	
103	E. Oxygen therapy in critical illness :
104	Cardiopulmonary resuscitation
105	Major trauma
106	Sepsis
107	Shock
108	Drowning
109	Anaphylaxis
110	Pulmonary haemorrhage
111	Seizures
112	Major head injury
113	Carbon monoxide poisoning
114	
115	F. Oxygen therapy for specific conditions that frequently require oxygen therapy:
116	Asthma
117	Pneumonia
118	Lung cancer
119	Pulmonary fibrosis and other interstitial lung disease,
120	Pneumothorax
121	Pleural effusion
122	Pulmonary embolism
123	Acute heart failure
124	Anaemia
125	Sickle cell crisis
126	Myocardial infarction and acute coronary syndrome
127	Stroke

- 128 Suspected hyperventilation
- 129 Most poisonings
- 130 Paraquat and bleomycin poisoning
- 131 Metabolic and renal disorders
- 132
- 133 G. Patients at risk of hypercapnic respiratory failure
 - 134 COPD
 - 135 Cystic Fibrosis
 - 136 Musculoskeletal and neurological disorders
 - 137 Morbid obesity
 - 138 Non Invasive Ventilation for acidotic type 2 respiratory failure
 - 139
- 140 H. Oxygen use during pregnancy
- 141 J. Oxygen use in perioperative care and during procedures requiring sedation procedures
- 142 K. Oxygen use in Palliative Care
- 143 L. Use of mixtures of oxygen with other gases (Heliox and Entonox)
- 144 M. Use of nitrous oxide-oxygen mixtures (Entonox) for analgesia
- 145 N. Use of CPAP in the perioperative period and for pulmonary oedema
- 146 P. Patients with tracheostomy or laryngectomy
- 147 Q. Humidification of oxygen
- 148 R. Driving gas for nebulised treatments
- 149 S. Prescribing oxygen therapy
- 150 T. Monitoring an adjusting oxygen therapy
- 151 U. Weaning and discontinuation of oxygen therapy
- 152 V. Practical aspects of oxygen use in pre-hospital and hospital care and oxygen alert cards
- 153 W. Practical aspects of oxygen dispensing, documentation and monitoring
- 154 X. Training in oxygen prescribing and use

155		
156	Table 1	Critical illnesses requiring high levels of supplemental oxygen
157	Table 2	Serious illnesses requiring moderate level of supplemental oxygen if the patient is hypoxaemic
158		
159	Table 3	Conditions for which patients should be monitored closely but oxygen therapy is not required unless the patient is hypoxaemic
160		
161	Table 4	COPD and other conditions requiring controlled or low-dose oxygen therapy
162	Table 5	Hierarchy of Evidence
163	Table 6	Grading of recommendations
164		
165		
166	Chart 1	Oxygen prescription guidance for acutely hypoxaemic patients in hospital
167	Chart 2	Flow chart for oxygen administration on general wards in hospitals
168		

169 References

170

171

172 *Healthcare providers need to use clinical judgement, knowledge and expertise when deciding*

173 *whether it is appropriate to apply recommendations for the management of patients. The*

174 *recommendations cited here are a guide and may not be appropriate for use in all situations. The*

175 *guidance provided does not override the responsibility of healthcare professionals to make decisions*

176 *appropriate to the circumstances of each patient, in consultation with the patient and/or their*

177 *guardian or carer.*

178

179 **Executive summary of the Guideline**

180 **Philosophy of the Guideline**

181

182 • Oxygen is a treatment for hypoxaemia, not breathlessness. Oxygen has not been
183 proven to have any consistent effect on the sensation of breathlessness in non-
184 hypoxaemic patients.

185

186 • The essence of this guideline can be summarised simply as a requirement for oxygen to
187 be prescribed according to a target saturation range and for those who administer
188 oxygen therapy to monitor the patient and keep within the target saturation range.

189

190 • The Guideline recommends aiming to achieve normal or near-normal oxygen
191 saturation for all acutely ill patients apart from those at risk of hypercapnic respiratory
192 failure or those receiving terminal palliative care.

193

194 **1. Assessing patients**

195

196 • For critically ill patients, high concentration oxygen should be administered immediately
197 (Table 1 and Chart 1) and this should be recorded afterwards in the patient's health
198 record.

199

200 • The oxygen saturation should be checked by pulse oximetry in all breathless and acutely ill
201 patients, "the fifth vital sign", (supplemented by blood gases when necessary) and the
202 inspired oxygen concentration should be recorded on the observation chart with the
203 oximetry result. (The other vital signs are Pulse rate, Blood Pressure, Temperature and
204 Respiratory Rate).

205

206 • Pulse oximetry must be available in all locations where emergency oxygen is used. Clinical
207 assessment is recommended if the saturation falls by $\geq 3\%$ or below the target range for
208 the patient.

209

210 • All critically ill patients should be assessed and monitored using a recognised physiological
211 track and trigger system such as the national early warning score (NEWS).

212

213 **2. Oxygen prescription**

214 • Oxygen should be prescribed to achieve a target saturation of 94-98% for most acutely ill
215 patients or 88-92% or patient-specific target range for those at risk of hypercapnic
216 respiratory failure. (Tables 1-4).

217

218 • Best practice is to prescribe a target range for all hospital patients at the time of admission
219 so that appropriate oxygen therapy can be commenced in the event of unexpected
220 clinical deterioration with hypoxaemia and also to ensure that the oximetry section of
221 the early warning score can be scored appropriately.

222

223 • The target saturation should be written (or ringed) on the drug chart or entered in an
224 electronic prescribing system (Guidance on Chart 1).

225

226

227

228

229 **3. Oxygen administration**

- 230 • Oxygen should be administered by staff who are trained in oxygen administration.
- 231
- 232 • These staff should use appropriate devices and flow rates in order to achieve the target
- 233 saturation range (Chart 2).
- 234

235 **4. Monitoring and maintenance of target saturation**

- 236 • Oxygen saturation and delivery system (including flow rate) should be recorded on the
- 237 patient's monitoring chart.
- 238
- 239 • Oxygen delivery devices and flow rates should be adjusted to keep the oxygen
- 240 saturation in the target range. Clinical assessment is required if oxygen therapy needs to
- 241 be initiated or increased due to a falling saturation level.
- 242
- 243 • Oxygen should be prescribed and a signature should be entered on the drug chart on
- 244 each drug round.
- 245

246 **5. Weaning and discontinuation of oxygen therapy**

- 247 • Oxygen should be reduced in stable patients with satisfactory oxygen saturation.
- 248
- 249 • Oxygen should be discontinued once the patient can maintain saturation within the
- 250 target range breathing air but the prescription for a target range should be left in place
- 251 in case of future deterioration and to guide early warning scores.

252 **Key changes since the first edition of this Guideline published in 2008**

253

254 **Methodology:**

255 The evidence review methodology has changed from NICE methodology to the BTS NICE accredited
256 guideline production process which is based on SIGN methodology and adheres to AGREE
257 methodology (see section 1).

258 **Evidence levels and grade of recommendation:**

259 These are now in SIGN format (see section 1 and tables 5 and 6).

260

261 **Evidence base:**

262 The evidence base for the Guideline has been updated to August 2013 (and extended to mid-2014
263 for key references). None of the 2008 recommendations have been challenged by new evidence but
264 many of the existing recommendations are supported by new information. There have been many
265 observational studies but few randomised trials directly relevant to the Guideline since 2008.

266

267 **The remit of the Guideline has been extended.**

268 The new Guideline covers not just emergency oxygen use but most oxygen use in health care
269 settings. It also covers short-term oxygen use by health care workers outside of health care settings
270 but domiciliary oxygen use by patients is covered by the BTS Guideline for home oxygen use in adults
271 [2].

272

273 **The scope of the guideline has been widened:**

274 The present Guideline includes the following new topics and settings which have been requested by
275 Guideline users:

- 276 • Postoperative and perioperative care including patient-controlled analgesia
- 277 • Endoscopy and other procedures requiring sedation
- 278 • Palliative care settings including hospices
- 279 • Use of helium-oxygen mixtures (Heliox) and nitrous oxide/oxygen mixtures (Entonox)
- 280 • Use of CPAP (continuous positive airway pressure)
- 281 • Use of oxygen by health care professionals in patients' homes
- 282 • Use of oxygen by voluntary rescue organisations and other non-NHS first responders
- 283 • High flow nasal cannulae

284

285 **The structure and format of the Guideline has been changed since 2008:**

286 The 2008 Guideline was published as a self-contained document in Thorax [1]. Additional
287 educational materials and other resources including audit tools were made available on the British
288 Thoracic Society website. The new Guideline exists in two complementary formats.

- 289 • A concise guideline which contains recommendations and good practice points is published
290 in Thorax.
- 291 • The full guideline including evidence review, physiology overview, illustrations and
292 references is available on the British Thoracic Society website www.brit-thoracic.org.uk

293

294

295 **1. INTRODUCTION**

296
297 **1.1 Aim of the Guideline**

298 The key aim of this Guideline is to make emergency oxygen use safer, simpler and more effective.
299 Oxygen is probably the commonest drug used in the care of patients who present with medical
300 emergencies. Prior to the publication of the first British Thoracic Society Guideline for Emergency
301 Oxygen Use in Adult Patients in 2008 [1], ambulance teams and emergency department teams were
302 likely to give oxygen to virtually all breathless or seriously ill patients and also to a large number of
303 non-hypoxaemic patients with conditions such as ischaemic heart disease or stroke based on custom
304 and practice. About 34% of UK ambulance journeys in 2007 involved oxygen use [3]. This translated
305 to about two million instances of emergency oxygen use per annum by all UK ambulance services,
306 with further use in patients' homes, GP surgeries and in hospitals. Audits of oxygen use and oxygen
307 prescription have shown consistently poor performance in many countries and most clinicians who
308 deal with medical emergencies have encountered adverse incidents and occasional deaths due to
309 underuse and overuse of oxygen [4-10].

310
311 Historically, oxygen has been administered for three main indications of which only one is evidence-
312 based. First, oxygen is given to correct hypoxaemia because severe hypoxaemia is clearly harmful to
313 the human body. Second, oxygen has been administered to ill patients in case they might become
314 hypoxaemic. Recent evidence suggests that, if impaired gas exchange does actually develop, this
315 practice may actually place patients at increased risk (see online section 3). Third, a very high
316 proportion of medical oxygen was administered because most clinicians believed, prior to 2008, that
317 oxygen can alleviate breathlessness in most circumstances. However, there is no good evidence that
318 oxygen relieves breathlessness in non-hypoxaemic patients. There is evidence of lack of
319 effectiveness or minimal effectiveness in mildly hypoxaemic breathless patients with chronic
320 obstructive pulmonary disease (COPD) and advanced cancer (see online section 6 and section
321 8.11.4).

322
323 Against this background, the Standards of Care Committee of the British Thoracic Society (BTS)
324 established a working party in association with 21 other societies to produce an evidence-based
325 guideline for emergency oxygen use in the UK. This led to the production of the 2008 British Thoracic
326 Society Guideline for emergency oxygen use in adult patients which was the world's first guideline
327 for emergency oxygen therapy [1]. This guideline has been implemented throughout the UK and in
328 many other countries leading to over 200 citations in the medical literature up to the end of 2013.

329
330 The purpose of the 2015 update to the guideline is to strengthen the evidence-base of the previous
331 guideline based on revised methodology (which meets criteria contained in the AGREE Instrument)
332 and to extend the evidence base to the end of 2013 and extended to mid 2014 for key references
333 [11]. Additionally, the remit of the 2008 Guideline has been broadened to cover several new aspects
334 of oxygen use and a broader range of locations where oxygen might be used.

335
336 **1.2 Intended users of the guideline and target patient populations**

337 This guideline is mainly intended for use by all healthcare professionals who may be involved in
338 emergency oxygen use. This will include ambulance staff, first responders, paramedics, doctors,
339 nurses, midwives, physiotherapists, pharmacists and all other healthcare professionals who may deal
340 with ill or breathless patients. Advice is also provided for first responders belonging to voluntary
341 organizations or other non-NHS bodies. Information based on this guideline is available on the BTS
342 website for use in the following situations:

- 343 • Hospital use
344 • Primary care use

- 345 • Ambulance use (supplemented by Ambulance service Guidance based on this guideline [12].
346 • Use by nursing staff and allied health professions

347 These abbreviated versions of the guideline contain the key recommendations and tables and charts
348 that are relevant to the particular situation. The “mini-guidelines” can be downloaded by health
349 care trusts for use on trust intranets and to produce paper versions of the guideline for key staff.

350

351 **1.3 Areas covered by this guideline**

352 The guideline addresses the use of oxygen in three main categories of adult patients in the pre-
353 hospital and hospital setting and in other settings such as palliative care:

- 354 • Critically ill or hypoxaemic patients.
355 • Patients at risk of hypoxaemia.
356 • Non-hypoxaemic patients who may benefit from oxygen (e.g., carbon monoxide poisoning).

357

358 **1.4 Areas not covered by this guideline**

- 359 • Oxygen use in paediatrics: the present guideline applies only to subjects aged >16 years.
360 • Oxygen use for high altitude activities.
361 • Oxygen use during air travel.
362 • Underwater diving and diving accidents.
363 • Oxygen use in animal experiments.
364 • Oxygen use in high-dependency units.
365 • Oxygen use in intensive care units.
366 • Inter-hospital level 3 transfers.
367 • Hyperbaric oxygen.
368 • Respiratory support techniques including tracheal intubation, invasive ventilation and non-
369 invasive ventilation (NIV) [Continuous Positive Airway Pressure (CPAP) is included]
370 • Self-initiated use of oxygen by patients who have home oxygen for any reason.
371 • Ongoing care of hypoxaemic patients at home.

372

373 **1.5 Limitations of the guideline**

374 This guideline is based on the best available evidence concerning oxygen therapy. However, a
375 guideline can never be a substitute for clinical judgement in individual cases. There may be cases
376 where it is appropriate for clinicians to act outwith the advice contained in this guideline because of
377 the needs of individual patients, especially those with complex or interacting disease states.
378 Furthermore, the responsibility for the care of individual patients rests with the clinician in charge of
379 the patient’s care and the advice offered in this guideline must, of necessity, be of a general nature
380 and should not be relied upon as the only source of advice in the treatment of individual patients. In
381 particular, this guideline gives very little advice about the management of the many medical
382 conditions that may cause hypoxaemia (apart from the specific issue of managing the patients’
383 hypoxaemia). Readers are referred to other guidelines for advice on the management of specific
384 conditions such as COPD, pneumonia, heart failure, etc. Some of these disease-specific guidelines
385 may suggest slightly different approaches to emergency oxygen therapy whereas the present
386 guideline aims to provide simple all-embracing advice about oxygen therapy.

387

388
389
390
391
392
393
394
395
396
397
398
399
400
401
402
403
404
405
406
407
408

2. METHODOLOGY OF GUIDELINE PRODUCTION

2.1 Methodology

The methodology for the 2015 Guideline is described in detail in the full Guideline which is available on the BTS website at: www.brit-thoracic.org.uk The 2008 Guideline used NICE methodology but the 2015 Guideline follows the BTS Guideline Production Manual 2014 which adheres to the AGREE instrument [11 13]. Significant new areas in the 2015 Guideline include the use of oxygen during procedures requiring sedation, the non-emergency use of oxygen in health care settings and the use of CPAP, Heliox and oxygen-nitrous oxide mixtures.

2.2 Planned review and updating of the guideline

2015 guideline will be reviewed by the BTS and by the endorsing organisations within 5 years from publication.

2.3 Declarations of interest

All members of the Guideline Group made declarations of interest in line with the BTS Policy and further details can be obtained on request from BTS.

409 **3. SUMMARY OF GUIDELINE RECOMMENDATIONS AND GOOD PRACTICE POINTS**

410

411 **A Achieving desirable oxygen saturation ranges in acute illness (see online sections 6 and**
412 **8 and Charts 1-2)**

413

414 A1. This guideline recommends aiming to achieve a normal or near-normal oxygen saturation for all
415 acutely ill patients apart from those at risk of hypercapnic respiratory failure. [Grade D]

416

417 A2. The recommended target saturation range for acutely ill patients not at risk of hypercapnic
418 respiratory failure is 94–98%. [Grade D]

419

420 A3. For most patients with known COPD or other known risk factors for hypercapnic respiratory
421 failure (e.g. morbid obesity, cystic fibrosis, chest wall deformities or neuromuscular disorders), a
422 target saturation range of 88–92% is suggested pending the availability of blood gas results. [Grade A
423 for COPD, Grade D for other conditions]

424

425 A4. Most non-hypoxaemic breathless patients do not benefit from oxygen therapy, but a sudden
426 reduction of $\geq 3\%$ in a patient's oxygen saturation within the target saturation range should prompt
427 fuller assessment of the patient (and the oximeter signal) because this may be the first evidence of
428 an acute illness. [Grade D]

429

430 A5. Because oxygenation is reduced in the supine position, fully conscious hypoxaemic patients
431 should ideally be allowed to maintain the most upright posture possible (or the most comfortable
432 posture for the patient) unless there are good reasons to immobilise the patient (e.g., skeletal or
433 spinal trauma). [Grade D]

434

435

436

437 **B Clinical and laboratory assessment of hypoxaemia and hypercapnia (see online section 7)**

438 B1. Fully trained clinicians should assess all acutely ill patients by measuring pulse rate, blood
439 pressure, temperature, respiratory rate and assessing circulating blood volume and anaemia. Expert
440 assistance from specialists in intensive care or from other disciplines should be sought at an early
441 stage if patients are thought to have major life-threatening illnesses and clinicians should be
442 prepared to call for assistance when necessary including a call for a 999 ambulance in pre-hospital
443 care or a call for the resuscitation team or ICU outreach team in hospital care. [Grade D]

444

445 B2. Initial clinical assessment and subsequent monitoring of acutely unwell patients should include
446 the use of a recognised physiological “track and trigger” system, such as the national early warning
447 score (NEWS) which may trigger clinical review due to hypoxaemia, need for supplementary oxygen
448 or for other reasons. [Grade D]

449

450 B3. Oxygen saturation, “the fifth vital sign”, should be checked by trained staff using pulse oximetry
451 in all breathless and acutely ill patients (supplemented by blood gases when necessary) and the
452 inspired oxygen device and flow rate should be recorded on the observation chart with the oximetry
453 result. [Grade D]

454

455 B4. The presence of a normal SpO₂ does not always negate the need for blood gas measurements
456 because pulse oximetry will be normal in a patient with normal oxygen tension but abnormal blood
457 pH or PCO₂ or with a low blood oxygen content due to anaemia. Blood gases and full blood count

458 tests are therefore required as early as possible in all situations where these measurements may
459 affect patient outcomes. [Grade D]

460

461 **Good Practice Points for clinical assessment of patients with suspected hypoxaemia**

- 462 • The medical history should be taken when possible in an acutely breathless patient and may
463 point to the diagnosis of a particular acute illness such as pneumonia or pulmonary
464 embolism or an exacerbation of a chronic condition such as COPD, asthma or heart failure.
- 465 • Do not discontinue oxygen therapy to obtain an oximetry measurement on room air in
466 patients who clearly require oxygen therapy.
- 467 • Physical examination may provide evidence of a specific diagnosis such as heart failure or a
468 large pleural effusion, but it is common for the cause of breathlessness to remain
469 undiagnosed until the results of tests such as chest radiographs are available.
- 470 • Patients with severe hypoxaemia may present with a non-respiratory manifestation such as
471 confusion or agitation rather than breathlessness and cyanosis is a difficult physical sign to
472 record confidently (especially in poor light or with an anaemic or plethoric patient).
- 473 • Tachycardia and tachypnoea are more common than a physical finding of cyanosis in
474 hypoxaemic patients.
- 475 • Appropriate changes should be made to any “track and trigger” system used to allow for a
476 lower target range in patients at risk of type 2 respiratory failure. These patients should
477 score no EWS points for saturation if within their target range and they should score points if
478 the oxygen saturation falls below the target range or if the saturation rises above the target
479 range whilst breathing oxygen.

480

481 **C Arterial and arteriased capillary blood gases (see online sections 7.1.3 and 8.4 and 8.5)**

482 C1. For critically ill patients or those with shock or hypotension (systolic blood pressure <90 mm Hg),
483 the initial blood gas measurement should be obtained from an arterial sample. For most patients
484 who require blood gas sampling, either arterial blood gases or arteriased earlobe blood gases may
485 be used to obtain an accurate measure of pH and PCO₂. However, the PaO₂ is less accurate in
486 earlobe blood gas samples (it underestimates the oxygen tension by 0.5– 1 kPa) so oximetry should
487 be monitored carefully if earlobe blood gas specimens are used and a repeat arterial specimen
488 should be taken if there is any concern about the accuracy of a capillary sample. [Grade D]

489

490 C2. Local anaesthesia should be used for all arterial blood gas specimens except in emergencies or if
491 the patient is unconscious or anaesthetised. [Grade A]

492

493 C3. Blood gases should be checked in the following situations:

494 – All critically ill patients.

495 – Unexpected or inappropriate fall in SpO₂ below 94% in patients breathing air or oxygen) or any
496 patient requiring oxygen to achieve the above target range. (Allowance should be made for transient
497 dips in saturation to 90% or less in normal subjects during sleep). [Grade D]

498 – Deteriorating oxygen saturation (fall of ≥3%.) or increasing breathlessness in a patient with
499 previously stable chronic hypoxaemia (e.g. severe COPD). [Grade D]

500 – Most previously stable patients who deteriorate clinically and require increased FiO₂ to maintain a
501 constant oxygen saturation. [Grade D]

502 – Any patient with risk factors for hypercapnic respiratory failure who develops acute
503 breathlessness, deteriorating oxygen saturation, drowsiness or other symptoms of carbon dioxide
504 retention. [Grade D]

505 – Patients with breathlessness who are thought to be at risk of metabolic conditions such as diabetic
506 ketoacidosis or metabolic acidosis due to renal failure. [Grade D]

507 – Any other evidence from the patient’s medical condition that would indicate that blood gas results
508 would be useful in the patient’s management (e.g. an unexpected change in “track and trigger”

509 systems such as a sudden rise of several units in the NEWS score or an unexpected fall in oxygen
510 saturation of 3% or more, even if within the target range). [Grade D]

511

512 **Good Practice Point: Patients requiring increased concentration of oxygen**

- 513 • The requirement for an increased concentration of oxygen is an indication for urgent clinical
514 reassessment of the patient (and repeat blood gas measurements in most instances, see
515 recommendations V13 and V18 for exceptions).

516

517

518 **D Initial oxygen therapy; initial choice of equipment for patients who do not have critical**
519 **illness (see charts 1-2 and table 2 and online section 8.9 and section 10).** *Initial oxygen*
520 *therapy in critical illness is covered in the next section*

521

522 D1. For acutely breathless patients not at risk of hypercapnic respiratory failure who have
523 saturations below 85%, treatment should be commenced with a reservoir mask at 15 l/min in the
524 first instance. The oxygen concentration can be adjusted downwards (using nasal cannulae or a
525 simple face mask) to maintain a target saturation of 94–98% once the patient has stabilised. [Grade
526 D]

527

528 D2. In other cases of acute hypoxaemia without critical illness or risk factors for hypercapnic
529 respiratory failure, treatment should be commenced with nasal cannulae (or a simple face mask if
530 cannulae are not tolerated or not effective) with the flow rate adjusted to achieve a saturation of
531 94–98%. [Grade D]

532

533 D3. If medium-concentration therapy with nasal cannulae or a simple face mask does not achieve
534 the desired saturation, change to a reservoir mask and seek senior or specialist advice. [Grade D]

535

536 **Good Practice Point**

- 537 • High flow nasal oxygen should be considered as an alternative to reservoir mask treatment
538 in patients with acute hypoxaemic (type 1) respiratory failure.

539

540 *For initial management of patients at risk of hypercapnic respiratory failure, see recommendations
541 G1 and G2

542

543

544

545

- 546 **E Oxygen therapy in critical illness (see table 1 and online section 8.10)**
547 E1. Use the highest feasible inspired oxygen for ventilation during **cardiopulmonary resuscitation**.
548 Once spontaneous circulation has returned, aim for a target saturation range of 94-98% and take an
549 arterial blood gas sample to guide on-going oxygen therapy. [Grade D]
550
551 E2. In **critical illness, including major trauma, sepsis**, shock and anaphylaxis, initiate treatment with
552 a reservoir mask at 15 l/min and aim at a saturation range of 94–98%. This advice also applies to
553 patients with critical illness who have risk factors for hypercapnia pending the results of blood gas
554 measurements and expert assessment. In patients with spontaneous circulation and a reliable
555 oximetry reading it may be possible to maintain a saturation of 94-98% using lower concentrations
556 of oxygen. Grade D
557
558 E3. In cases of **drowning**, aim at an oxygen saturation of 94–98% once spontaneous circulation is
559 restored. [Grade D]
560
561 E4. In patients with **acute seizures due to epilepsy or other causes**, high concentration oxygen
562 should be administered until a satisfactory oximetry measurement can be obtained and clinicians
563 should then aim an oxygen saturation of 94–98% or 88–92% if the patient is at risk of hypercapnic
564 respiratory failure. [Grade D]
565
566 E5. In cases of **major head injury**, aim at an oxygen saturation of 94–98%. Initial treatment should
567 involve high concentration oxygen from a reservoir mask at 15 l/min pending availability of
568 satisfactory blood gas measurements or until the airway is secured by intubation. [Grade D]
569
570 E6. In cases of **carbon monoxide poisoning**, an apparently “normal” oximetry reading may be
571 produced by carboxyhaemoglobin, so aim at an oxygen saturation of 100% and use a reservoir mask
572 at 15 l/min irrespective of the oximeter reading and PaO₂. [Grade D]
573

Table 1 Critical illnesses requiring high levels of supplemental oxygen Section 8.10

<p>The initial oxygen therapy is a reservoir mask at 15 l/min pending the availability of reliable oximetry readings.</p> <p>For patients with spontaneous circulation and a reliable oximetry reading, it may quickly become possible to reduce the oxygen dose whilst maintaining a target saturation range of 94-98%.</p> <p>If oximetry is unavailable, continue to use a reservoir mask until definitive treatment is available.</p> <p>Patients with COPD and other risk factors for hypercapnia who develop critical illness should have the same initial target saturations as other critically ill patients pending the results of blood gas results after which these patients may need controlled oxygen therapy or supported ventilation if there is severe hypoxaemia and/or hypercapnia with respiratory acidosis.</p>		
	Additional Comments	Recommendations
Cardiac arrest or resuscitation	<p>Refer to resuscitation guidelines for choice of delivery device during active resuscitation.</p> <p>Give highest possible inspired oxygen concentration during CPR until spontaneous circulation has been restored.</p>	Recommendation E1
Shock, sepsis, major trauma, drowning, anaphylaxis, major pulmonary haemorrhage, status epilepticus	Also give specific treatment for the underlying condition	Recommendations E2-E4
Major Head Injury	Early tracheal intubation and ventilation if comatose.	Recommendation E5
Carbon Monoxide Poisoning	<p>Give as much oxygen as possible using a bag-valve mask or reservoir mask. Check carboxyhaemoglobin levels.</p> <p>A normal or high oximetry reading should be disregarded because saturation monitors cannot differentiate between carboxyhaemoglobin and oxyhaemoglobin, owing to their similar absorbances.</p> <p>The blood gas PaO₂ will also be normal in these cases (despite the presence of tissue hypoxia).</p>	Recommendation E6

COPD Chronic obstructive pulmonary disease;
PaO₂ arterial oxygen tension

576 **F Oxygen therapy for specific conditions that frequently require oxygen therapy**
577 **(see Tables 2 and 3 and online section 8.11 and 8.13)**

578
579 **Respiratory conditions**

- 580
581 F1. In **acute asthma**, aim at an oxygen saturation of 94–98%. [Grade D]
582
583 F2. In cases of **pneumonia** who are not at risk of hypercapnic respiratory failure, aim at an oxygen
584 saturation of 94–98%. [Grade D]
585
586 F3. In acute **breathlessness due to lung cancer**, aim at an oxygen saturation of 94–98% unless there
587 is coexisting COPD. See also ‘Oxygen use in Palliative Care’. [Grade D]
588
589 F4. In **acute deterioration of pulmonary fibrosis or other interstitial lung diseases**, aim at an
590 oxygen saturation of 94–98% or the highest possible if these targets cannot be achieved. [Grade D]
591
592 F5. In most cases of **pneumothorax**, aim at an oxygen saturation of 94–98% if the patient is not at
593 risk of hypercapnic respiratory failure. [Grade D]
594
595 F6. In patients with pneumothorax having hospital observation without drainage, the use of high
596 concentration oxygen (15 l/min flow rate via reservoir mask) is recommended unless the patient is at
597 risk of hypercapnic respiratory failure. [Grade D]
598
599 F7. In **pleural effusion**, aim at an oxygen saturation of 94–98% (or 88–92% if the patient is at risk of
600 hypercapnic respiratory failure). [Grade D]
601
602 F8. In **pulmonary embolism**, aim at an oxygen saturation of 94–98% (or 88–92% if the patient is at
603 risk of hypercapnic respiratory failure). [Grade D]
604

605 **Non-respiratory conditions**

- 606
607 F9. In **acute heart failure**, aim at an oxygen saturation of 94–98% (or 88–92% if the patient is at risk
608 of hypercapnic respiratory failure). [Grade D]
609
610 F10. Consider treatment with continuous positive airway pressure to relieve symptoms of heart
611 failure if there is hypoxaemia or treatment with non-invasive ventilation if there is co-existent
612 hypercapnia with acidosis. [Grade B]
613
614 F11. In **anaemia**, aim at an oxygen saturation of 94–98% or 88–92% if the patient is at risk of
615 hypercapnic respiratory failure. [Grade D]
616
617 **Good Practice Point:**
618 Correction of anaemia by blood transfusion should be based on national guidelines.
619 F12. In **sickle cell crisis** and acute chest syndrome, aim for an oxygen saturation of 94–98% or aim at
620 the saturation level that is usual for the individual patient. [Grade D]
621

622 **Good Practice Point regarding sickle cell crisis**

623 Arterial or arterialised capillary blood gases should be sampled if there is any doubt about the
624 reliability of oximetry during a sickle cell crisis.

625

626 F13. In **myocardial infarction and acute coronary syndromes**, aim at an oxygen saturation of 94–
627 98% or 88–92% if the patient is at risk of hypercapnic respiratory failure. [Grade D]

628

629 F14. High concentrations of oxygen should be avoided in **stroke patients**, unless required to
630 maintain normal oxygen saturation. Aim at an oxygen saturation of 94–98% or 88–92% if the patient
631 is at risk of hypercapnic respiratory failure. [Grade D]

632

633 **Good Practice Points regarding stroke management**

- 634 • Oxygen saturation should be monitored at least every 4 hours throughout the day and night
635 in patients with acute stroke and all episodes of hypoxaemia treated.
- 636 • Patients with hypoxaemia post stroke require medical review to establish and treat the
637 cause.
- 638 • Oxygen should only be given once the airway has been cleared and at the lowest
639 concentration necessary to achieve an oxygen saturation of 94–98% or 88–92% if the patient
640 is at risk of hypercapnic respiratory failure.
- 641 • Oxygen should be given via nasal cannulae, unless there are clear indications for a different
642 oxygen delivery system.
- 643 • Patients with stroke and cardio-respiratory comorbidities should be positioned as upright as
644 possible, in a chair if possible. **(see recommendation A5)**
- 645 • Patients with a reduced level of consciousness after stroke should be nursed in the recovery
646 position.

647

648 **Suspected hyperventilation**

649 **Good practice points regarding patients with suspected hyperventilation**

- 650 • Organic illness must be excluded before making a diagnosis of hyperventilation.
- 651 • Patients with a definite diagnosis of hyperventilation should have their oxygen saturation
652 monitored. Those with normal or high SpO₂ do not require oxygen therapy.
- 653 • Rebreathing from a paper bag can be dangerous and is NOT advised as a treatment for
654 hyperventilation.

655

656 F15. In **most poisonings**, aim at an oxygen saturation of 94–98% unless the patient is at risk of
657 hypercapnic respiratory failure. [Grade D]

658

659 F16. In **poisoning by paraquat and bleomycin**, give oxygen only if the saturation falls below 85%
660 and reduce or stop oxygen therapy if the saturation rises above 88% [Grade D]

661

662 F17. In most **metabolic and renal disorders**, aim at an oxygen saturation of 94–98% unless the
663 patient is at risk of hypercapnic respiratory failure. [Grade D]

664

665

Table 2 Serious illnesses requiring moderate levels of supplemental oxygen if the patient is hypoxaemic Section 8.11

<p>The initial oxygen therapy is nasal cannulae at 2-6 l/min (preferably), or simple face mask at 5-10 l/min unless stated otherwise.</p> <p>For patients not at risk of hypercapnic respiratory failure who have saturation below 85%, treatment should be commenced with a reservoir mask at 15 l/min and the recommended initial oxygen saturation target range is 94-98%. If oximetry is not available, give oxygen as above until oximetry or blood gas results are available. Change to reservoir mask if the desired saturation range cannot be maintained with nasal cannulae or simple face mask (and ensure that the patient is assessed by senior medical staff).</p> <p>If these patients have co-existing COPD or other risk factors for hypercapnic respiratory failure, aim at a saturation of 88-92% pending blood gas results but adjust to 94-98% if the PaCO₂ is normal (unless there is a history of previous hypercapnic respiratory failure requiring NIV or IPPV) and recheck blood gases after 30-60 minutes.</p>		
	Additional Comments	Recommendation
Acute hypoxaemia (cause not yet diagnosed)	Reservoir mask at 15 l/min if initial SpO ₂ below 85%, otherwise nasal cannulae or simple face mask. <i>Patients requiring reservoir mask therapy need urgent clinical assessment by senior staff.</i>	Recommendations D1-D3
Acute Asthma Pneumonia Lung Cancer		Recommendations F1-F3
Deterioration of lung fibrosis or other interstitial lung disease	Reservoir mask at 15 l/min if initial SpO ₂ below 85%, otherwise nasal cannulae or simple face mask	Recommendation F4
Pneumothorax	Needs aspiration or drainage if the patient is hypoxaemic. Most patients with pneumothorax are not hypoxaemic and do not require oxygen therapy. Use a reservoir mask at 15 l/min if admitted for observation without chest drainage. Aim at 100% saturation to accelerate clearance of pneumothorax.	Recommendations F5-F6
Pleural Effusions	Most patients with pleural effusions are not hypoxaemic. If hypoxaemic, treat by draining the effusion as well as giving oxygen therapy.	Recommendation F7
Pulmonary Embolism	Most patients with minor pulmonary embolism are not hypoxaemic and do not require oxygen therapy.	Recommendation F8
Acute Heart Failure	Consider CPAP or NIV in cases of pulmonary oedema.	Recommendations F9-F10
Severe Anaemia	The main issue is to correct the anaemia. Most anaemic patients do not require oxygen therapy.	Recommendations F11-12
Post-operative Breathlessness	Management depends on underlying cause.	Recommendation J1

COPD-chronic obstructive pulmonary disease; CPAP- continuous positive airway pressure; IPPV- intermittent positive pressure ventilation; NIV – non-invasive ventilation; PaCO₂ arterial carbon dioxide tension, SpO₂-arterial oxygen saturation measured by pulse oximetry

666
667

Table 3 Conditions for which patients should be monitored closely but oxygen therapy is not required unless the patient is hypoxaemic Section 8.13

<p>If hypoxaemic, the initial oxygen therapy is nasal cannulae at 2-6 l/min or simple face mask at 5-10 l/min unless saturation is below 85% (use reservoir mask) or if at risk from hypercapnia. (see below)</p> <p>The recommended initial target saturation range, unless stated otherwise, is 94-98%. If oximetry is not available, give oxygen as above until oximetry or blood gas results are available.</p> <p>If patients have COPD or other risk factors for hypercapnic respiratory failure, aim at a saturation of 88-92% pending blood gas results but adjust to 94-98% if the PaCO₂ is normal (unless there is a history of respiratory failure requiring NIV or IPPV) and re-check blood gases after 30-60 minutes.</p>		
	Additional Comments	Recommendations
Myocardial infarction and Acute Coronary Syndromes	Most patients with acute coronary artery syndromes are not hypoxaemic and the benefits/harms of oxygen therapy are unknown in such cases. Un-necessary use of high concentration oxygen may increase infarct size.	<i>Recommendation F13</i>
Stroke	Most stroke patients are not hypoxaemic. Oxygen therapy may be harmful for non-hypoxaemic patients with mild-moderate strokes.	<i>Recommendation F14</i>
Hyperventilation or dysfunctional breathing	Exclude organic illness. Patients with pure hyperventilation due to anxiety or panic attacks are unlikely to require oxygen therapy. Re-breathing from a paper bag may cause hypoxaemia and is not recommended.	<i>See section 8.13.3</i>
Most poisonings and drug overdoses <i>(See Table 1 for Carbon Monoxide poisoning)</i>	Hypoxaemia is more likely with respiratory depressant drugs, give antidote if available. e.g. <i>Naloxone for opiate poisoning</i> Check blood gases to exclude hypercapnia if a respiratory depressant drug has been taken. Avoid high blood oxygen levels in cases of acid aspiration as there is theoretical evidence that oxygen may be harmful in this condition. Monitor all potentially serious cases of poisoning in a Level 2 or Level 3 environment (High Dependency Unit or Intensive Care Unit)	<i>Recommendation F15</i>
Poisoning with Paraquat or Bleomycin	Patients with Paraquat poisoning or Bleomycin lung injury may be harmed by supplemental oxygen. Avoid oxygen unless the patient is hypoxaemic. Target saturation is 85-88%.	<i>Recommendation F16</i>
Metabolic & Renal disorders	Most do not need oxygen (Tachypnoea may be due to acidosis in these patients)	<i>Recommendation F17</i>
Acute and sub-acute neurological and muscular conditions producing muscle weakness	These patients may require ventilatory support and they need careful monitoring which includes spirometry. If the patient's oxygen level falls below the target saturation, they need urgent blood gas measurements and are likely to need ventilatory support.	<i>Recommendation G4</i>
Pregnancy and Obstetric Emergencies	Oxygen therapy may be harmful to the foetus if the mother is not hypoxaemic.	<i>Recommendations H1-H4</i>

COPD-chronic obstructive pulmonary disease; IPPV- intermittent positive pressure ventilation; NIV –non-invasive ventilation; PaCO₂ arterial carbon dioxide tension, SpO₂-arterial oxygen saturation measured by pulse oximetry

669 **G Patients at risk of hypercapnic respiratory failure. See Table 4 and online Section 8.12**

670

671 G1 (also A3). For most patients with known COPD or other known risk factors for hypercapnic
672 respiratory failure (e.g. morbid obesity, cystic fibrosis, chest wall deformities or neuromuscular
673 disorders), a target saturation range of 88–92% is suggested pending the availability of blood gas
674 results. [Grade A for COPD, Grade D for other conditions]

675

676 G2. Some patients with COPD and other conditions are vulnerable to repeated episodes of
677 hypercapnic respiratory failure. In these cases it is recommended that treatment should be based on
678 the results of previous blood gas estimations during acute exacerbations. For patients with prior
679 hypercapnic failure (requiring non-invasive ventilation or intermittent positive pressure ventilation)
680 who do not have an alert card, it is recommended that treatment should be commenced using either
681 nasal cannulae or a 24% Venturi mask at 2–4 l/min (or a 28% Venturi mask at 4 l/min if a 24% mask
682 is not available) with an initial target saturation of 88–92% pending urgent blood gas results. These
683 patients should be treated as a high priority by emergency services and the oxygen concentration
684 should be reduced if the saturation exceeds 92%. [Grade D]

685

686 **Good practice points for COPD and other conditions that may cause hypercapnic respiratory**
687 **failure:**

688 **Diagnosis of COPD or suspected exacerbation of COPD**

- 689 • If the diagnosis is unknown, patients over 50 years of age who are long-term smokers with a
690 history of chronic breathlessness on minor exertion such as walking on level ground and no
691 other known cause of breathlessness should be treated as if having COPD for the purposes
692 of this guideline.
- 693 • Measurement of spirometry may confirm (or exclude) a diagnosis of airflow obstruction and
694 the FEV1 level is a useful indicator of disease severity in COPD. Spirometry should be
695 measured at least once during hospital admissions for suspected COPD.

696

697 **Immediate management of patients with known or suspected COPD**

- 698 • Patients with a significant likelihood of severe COPD or other illness that may cause
699 hypercapnic respiratory failure should be triaged as very urgent on arrival in hospital
700 emergency departments and blood gases should be measured on arrival in hospital.
- 701 • Prior to availability of blood gas measurements, use a 28% Venturi mask at 4 l/min or 24%
702 Venturi mask or nasal cannulae at 2 l/min and aim for an oxygen saturation of 88–92%
- 703 • Patients with a respiratory rate >30 breaths/min should have the flow rate set to 50% above
704 the minimum flow rate specified for the Venturi mask packaging to compensate for the
705 patient's increased inspiratory flow. Increasing the oxygen flow rate into a Venturi mask
706 does not increase the concentration of oxygen which is delivered.
- 707 • If the saturation remains below 88% in pre-hospital care despite a 28% Venturi mask, change
708 to nasal cannulae at 2–6 l/min or a simple face mask at 5 l/min with target saturation of 88–
709 92% and alert the A&E department that the patient is to be treated as a high priority.

710

711 **Initial hospital management of patients with exacerbation of COPD**

- 712 • Patients with exacerbations of COPD need careful monitoring for hypercapnic (type 2)
713 respiratory failure with respiratory acidosis which may develop in the course of a hospital
714 admission even if the initial blood gases were satisfactory.
- 715 • The risk of respiratory acidosis in patients with hypercapnic respiratory failure is increased if
716 the arterial oxygen tension is above 10.0 kPa due to previous excessive oxygen use.
- 717 • If following blood gas measurements the pH and PCO₂ are normal, aim for an oxygen
718 saturation of 94–98% unless there is a history of previous hypercapnic respiratory failure
719 requiring non-invasive ventilation or intermittent positive pressure ventilation or if the

720 patient's usual oxygen saturation when clinically stable is below 94% (these patients should
721 have a target range of 88-92%). Blood gases should be repeated at 30-60 minutes to check
722 for rising PaCO₂ or falling pH.

- 723 • Recheck blood gases after 30–60 min (or if there is evidence of clinical deterioration) for all
724 patients with COPD or other risk factors for hypercapnic respiratory failure even if the initial
725 PaCO₂ measurement was normal.
- 726 • If the PaCO₂ is raised but pH is ≥ 7.35 ($[H^+] \leq 45$ nmol/l), the patient has probably got long-
727 standing hypercapnia; maintain target range of 88–92% for these patients. Blood gases
728 should be repeated at 30-60 minutes to check for rising PaCO₂ or falling pH.
- 729 • If the patient is hypercapnic (PaCO₂ >6 kPa or 45 mm Hg) and acidotic (pH <7.35 or $[H^+] >45$
730 nmol/l), consider non-invasive ventilation, especially if the acidosis has persisted for more
731 than 30 min despite appropriate therapy.
- 732 • For patients using Venturi masks, consider changing from Venturi mask to nasal cannulae
733 once the patient has stabilised.

734

735 **Good practice point.**

736 **Management of hypercapnia or respiratory acidosis due to excessive oxygen therapy**

- 737 • If a patient is suspected to have hypercapnia or respiratory acidosis due to excessive oxygen
738 therapy, the oxygen therapy should not be discontinued but should be stepped down to 28%
739 or 24% oxygen from a Venturi mask or 1-2 l/min via nasal cannulae depending on oxygen
740 saturation and subsequent blood gas results.

741

742

743 G3. Initial oxygen treatment of **cystic fibrosis exacerbations** should be similar to the initial oxygen
744 treatment of COPD exacerbations with target saturation 88-92% (see sections 8.12.1-8.12.2). [Grade
745 D]

746

747 G4. In the initial management of **musculoskeletal and neurological disorders** with acute respiratory
748 failure, aim at an oxygen saturation of 88–92% and measure blood gases to determine if non-
749 invasive ventilation will be required. [Grade D]

750

751 **Good practice point regarding patients with neurological disorders:**

752 Patients with respiratory failure due to acute, sub-acute or chronic neurological disorders or muscle
753 disease are likely to require non-invasive or invasive ventilator support rather than oxygen therapy.

754

755 G5. **Morbidly obese patients** (BMI > 40 Kg/m²), even without evidence of coexistent obstructive
756 sleep apnoea are at risk of hypoventilation and should be given titrated oxygen to maintain a target
757 saturation of 88-92% (Grade D)

758

759 G6. **Non-invasive ventilation** should be considered for hypercapnic patients with COPD at risk of
760 hypercapnic respiratory failure, cystic fibrosis, neuro-muscular disorders or morbid obesity if the pH
761 is <7.35 or $[H^+] >45$ nmol/l. [Grade D]

762

763

764

Table 4 COPD and other conditions requiring controlled or low-dose oxygen therapy
Section 8.12

<p>Prior to availability of blood gases, use a 28% Venturi mask at 4 l/min or nasal cannulae at 1-2 l/min and aim for an oxygen saturation of 88-92% for patients with risk factors for hypercapnia but no prior history of respiratory acidosis (Recommendation 4). Adjust target range to 94-98% if the PaCO₂ is normal (unless there is a history of previous NIV or IPPV) and recheck blood gases after 30-60 minutes.</p>		
	Additional Comments	Recommendations
COPD	May need lower range if acidotic or if known to be very sensitive to oxygen therapy. Ideally use "Alert cards" to guide therapy based on previous blood gas results. Increase flow by 50% if respiratory rate is above 30 breaths per minute.	Recommendations G1-G2 and Section 8.12.1
Exacerbation of Cystic Fibrosis	Admit to regional CF centre if possible, if not discuss with regional centre or manage according to protocol agreed with regional CF centre. Ideally use "Alert cards" to guide therapy. Increase flow by 50% if respiratory rate is above 30 breaths per minute.	Recommendations G1, G3, G6
Chronic Neuro-Muscular Disorders	May require ventilatory support. Risk of hypercapnic respiratory failure.	Recommendations G1, G4, G6
Morbid Obesity		Recommendations G1, G5, G6

CF-Cystic fibrosis; COPD -chronic obstructive pulmonary disease; CPAP- continuous positive airway pressure; IPPV- intermittent positive pressure ventilation; NIV –non-invasive ventilation; PaCO₂ arterial carbon dioxide tension, SpO₂-arterial oxygen saturation measured by pulse oximetry

765

Public consultation

766
767
768
769
770
771
772
773
774
775
776
777
778
779
780
781
782
783
784
785
786
787
788
789
790
791
792
793
794
795
796
797
798
799
800
801
802
803
804
805
806
807
808
809
810
811
812

H Oxygen use during pregnancy (see online section 8.14)

H1. Women who suffer from major trauma, sepsis or acute illness during pregnancy should receive the same oxygen therapy as any other seriously ill patients, with a target oxygen saturation of 94–98%. The same target range should be applied to women with hypoxaemia due to acute complications of pregnancy (e.g., collapse related to amniotic fluid embolus, eclampsia or antepartum or postpartum haemorrhage). [Grade D]

H2. Women with underlying hypoxaemic conditions (e.g., heart failure) should be given supplemental oxygen during labour to achieve an oxygen saturation of 94–98% unless they are at risk of hypercapnic respiratory failure (target range 88-92%). [Grade D]

H3. All women with evidence of hypoxaemia who are more than 20 weeks pregnant should be managed with left lateral tilt or manual displacement of the uterus to improve cardiac output and oxygen delivery. [Grade D]

H4. The use of oxygen supplementation during intrauterine fetal resuscitation during labour was widespread in the past but there is no evidence of benefit. There is weak evidence of harm to the foetus if supplemental oxygen is given for long periods during uncomplicated labour. Overall, the use of oxygen during labour is only required when there is evidence of maternal hypoxaemia (oxygen saturation less than 94%). [Grade D]

J Oxygen use in perioperative care and during procedures requiring sedation (see online sections 8.15-8.16)

J1. Hyperoxaemia is not recommended routinely in the perioperative and postoperative period to reduce the incidence of postoperative nausea and vomiting (Grade D)

J2. All procedures involving sedation warrant routine continuous monitoring of oxygen saturation via pulse oximetry prior and during the procedure, and in the recovery period, particularly fibre-optic bronchoscopy and upper GI endoscopy where a reduction in arterial oxygen saturation is common, particularly with concurrent use of sedation. [Grade C]

J3. Significant arterial oxygen desaturation ($SpO_2 < 90\%$ or or fall of 4% or more that is prolonged (>1 minute during endoscopy procedures) should be corrected by supplemental oxygen with the aim of achieving target oxygen saturations of 94-98%, or 88-92% in those at risk of hypercapnic respiratory failure. [Grade D]

J4. Complicated upper GI endoscopy or procedures in patients with cardiorespiratory comorbidity are especially likely to lead to hypoxaemia and may also lead to hypercapnia, especially if the patient is heavily sedated. It is recommended that blood gases should be measured if such patients should require prolonged oxygen administration. The routine administration of oxygen is not recommended as it may delay the recognition of respiratory failure. [Grade D]

J5. Constant clinical assessment of the patient is crucial at all stages of sedation procedures and monitoring of capnography or transcutaneous carbon dioxide levels may be a useful adjunct to identify early respiratory depression. [Grade D]

813 J6. During the recovery period after procedures requiring sedation, supplemental oxygen should be
814 titrated to achieve target saturations of 94-98% in most patients and 88-92% in those at risk of
815 hypercapnic respiratory failure (see 10.5.1). (Grade D)

816

817

818 **Good Practice Points related to oxygen use in perioperative care**

819 • A target saturation of 94-98% is recommended for most surgical patients except those at risk
820 of type 2 respiratory failure when a range of 88-92% should be achieved.

821 • Pulse oximetry monitoring is recommended for postoperative patients despite the lack of
822 evidence from randomised studies.

823 • Patients using **Patient Controlled Analgesia** should have two hourly oximetry observations
824 because of the risk of hypoxaemia. Oxygen should be administered to keep patients within
825 the appropriate target saturation range.

826 • A target saturation of 94-98% is advised in most patients having PCA except those at risk of
827 type 2 respiratory failure when a range of 88-92% should be achieved.

828 • There is conflicting evidence concerning the balance of potential benefits and risks of
829 **perioperative hyperoxaemia to reduce the risk of surgical site infection in elective surgery**
830 and there is no evidence for this practice in patients having emergency surgical procedures.
831 More trials are required for specific procedures and more information is required concerning
832 long-term mortality risks to cancer patients.

833 • Oxygen should not be used routinely for this indication and not used outside of clinical trials
834 except in specialised centres.

835

836 **K Oxygen use in Palliative Care (see online section 8.17)**

837 K1. Oxygen use in palliative care patients should be restricted to patients with severe hypoxaemia
838 or patients who report significant relief of breathlessness from oxygen; In non-hypoxaemic patients,
839 opioids and non-pharmacological measures should be tried before oxygen. (Grade B)

840

841 K2. In general, there is no role for the monitoring of oxygen saturation or PaO₂ in comfort-focused
842 terminal care at the end of life. If the patient appears comfortable, oxygen levels are irrelevant and
843 should not influence care. (Grade D)

844

845 **Good Practice Points related to oxygen use in palliative care**

846

847 Oxygen therapy for the symptomatic relief of dyspnoea in palliative care patients is more complex
848 than the simple correction of hypoxaemia. Consider the following issues:

849

850 • Consider early involvement of palliative care specialists.

851 • As dyspnoea is a multi-factorial sensation-a comprehensive assessment of contributing
852 factors (such as anxiety) should be carried out.

853 • Opioids are effective for the relief of breathlessness in palliative care patients.

854 • A trial of non-pharmacological delivery of air to help relieve dyspnoea (e.g. hand held fan) is
855 recommended prior to trial of oxygen.

856 • Oxygen use has to be tailored to the individual and a formal assessment made of its efficacy
857 for reducing breathlessness and improving quality of life for that person.

858 • Oxygen therapy should not be continued in the absence of benefit.

859

860 **L Mixtures of oxygen with other gases (Heliox and Entonox)**

861

862 **Use of helium-oxygen mixtures (Heliox) See online section 8.18**

863 L1. There is insufficient evidence to support the use of Heliox either as an inhaled gas or as the
864 driving gas for nebuliser therapy in adult patients with acute exacerbations of asthma or COPD
865 except as part of randomised clinical trials or in exceptional circumstances. (Grade D)

866

867 L2. Heliox use for asthma or COPD patients should be considered only in severe exacerbations in
868 patients who are not responding to standard treatment (and in COPD patients where there are
869 contra-indications to intubation). (Grade D)

870

871 L3. A therapeutic trial of Heliox is reasonable in patients with mechanical upper airway obstruction
872 or postoperative stridor. (Grade D)

873

874 **M Use of nitrous oxide/oxygen mixtures (Entonox) for analgesia. (See online section 9.11)**

875 M1. The use of Entonox gas mixture for analgesia should be avoided if possible in patients at risk of
876 type 2 respiratory failure. [Grade D]

877

878

879 **N CPAP and high flow nasal oxygen**

880 **Use of CPAP in the perioperative period and for pulmonary oedema (see online section**
881 **8.19)**

882 N1. Patients with diagnosed sleep disordered breathing established on CPAP undergoing surgery
883 should bring their machines with them and use them in the pre and post-operative period. If
884 adequate saturations are not achieved despite CPAP therapy then assess for worsening ventilation
885 with blood gases and oxygen should be entrained to achieve a saturation of 88-92% (Grade D)

886

887 N2. CPAP with entrained oxygen to maintain saturation 94-98% should be considered as an adjunctive
888 treatment to improve gas exchange in patients with cardiogenic pulmonary oedema who are not
889 responding to standard treatment in hospital care or in pre-hospital care. [Grade B]

890

891 **Good Practice Point, High Flow Nasal oxygen**

- 892 • High flow nasal oxygen should be considered as a potentially superior alternative to
893 reservoir mask treatment in patients with acute hypoxaemic (type 1) respiratory failure.

894

895 **P Patients with tracheostomy or laryngectomy (see section 10.3)**

896 P1. When oxygen is required by patients with prior tracheostomy or laryngectomy, a tracheostomy
897 mask (varying the flow as necessary) should achieve the desired oxygen saturation (tables 1–4). An
898 alternative delivery device, usually a T-piece device fitted directly to the tracheostomy tube, may be
899 necessary if the patient deteriorates. [Grade D]

900

901 **Q Humidification of oxygen (see section 10.2)**

902 Q1. Humidification is not required for the delivery of low-flow oxygen (mask or nasal cannulae) or
903 for the short-term use of high-flow oxygen. It is not therefore required in pre-hospital care. Pending
904 the results of clinical trials, it is reasonable to use humidified oxygen for patients who require high-
905 flow oxygen systems for more than 24 h or who report upper airway discomfort due to dryness.
906 [Grade D]

907

908 Q2. In the emergency situation, humidified oxygen use can be confined to patients with
909 tracheostomy or an artificial airway although these patients can be managed without humidification
910 for short periods of time (e.g., ambulance journeys). [Grade D]

911

912 Q3. Humidification may also be of benefit to patients with viscous secretions causing difficulty with
913 expectoration. This benefit can be achieved using nebulised normal saline. [Grade D]

914

915 Q4. Bubble bottles which allow a stream of oxygen to bubble through a container of water should
916 not be used because there is no evidence of a clinically significant benefit but there is a risk of
917 infection. [Grade D]

918

919 **Good practice points related to humidified oxygen therapy**

- 920 • Patients requiring high flow rates or longer term oxygen might benefit from a large volume
921 oxygen humidifier device, especially if sputum retention is a clinical problem.
- 922 • In the absence of an artificial airway the decision to humidify supplemental oxygen needs to
923 be made on an individual basis but this practice is not evidence-based.

924

925

926 **R Driving gas for nebulised treatments (see section 10.4)**

927 R1. For patients with asthma, nebulisers should be driven by piped oxygen or from an oxygen
928 cylinder fitted with a high-flow regulator capable of delivering a flow rate of >6 l/min. The patient
929 should be changed back to his/her usual oxygen mask or cannulae when nebuliser therapy is
930 complete. If the cylinder does not produce this flow rate, an air-driven nebuliser (with electrical
931 compressor) should be used with supplemental oxygen by nasal cannulae at 2–6 l/min to maintain
932 an appropriate oxygen saturation level. [Grade D]

933

934 R2. When nebulised bronchodilators are given to patients with hypercapnic acidosis, they should be
935 given using an ultrasonic nebuliser or else a jet nebuliser driven by compressed air and, if necessary,
936 supplementary oxygen should be given concurrently by nasal cannulae to maintain an oxygen
937 saturation of 88–92%. The same precautions should be applied to patients who are at risk of
938 hypercapnic respiratory failure prior to the availability of blood gas results and the oxygen saturation
939 should be monitored continuously during treatment. Once the nebulised treatment is completed for
940 patients at risk of hypercapnic respiratory failure, their previous targeted oxygen therapy should be
941 reinstated. [Grade D]

942

943 **Good practice point: Driving gas for nebulised treatment in ambulances**

- 944 • During treatment by ambulance staff oxygen-driven nebulisers should be used for patients
945 with asthma and may be used for patients with COPD in the absence of an air-driven
946 compressor system. If oxygen is used for patients with known COPD, its use should be
947 limited to 6 min. This will deliver most of the nebulised drug dose but limit the risk of
948 hypercapnic respiratory failure (section 10.4). Ambulance Services are encouraged to
949 explore the feasibility of introducing battery powered, air driven nebulisers or portable
950 ultrasonic nebulisers.

951

952

953 **S Prescribing, oxygen therapy (see online section 11)**

954 S1. Every healthcare facility should have a standard oxygen prescription document or, preferably, a
955 designated oxygen section on all drug prescribing cards or guided prescription of oxygen in
956 electronic prescribing systems. [Grade D]

957

958 S2. A prescription for oxygen should always be written, except in critical illness when it must be
959 started immediately and documented retrospectively. [Grade D]
960

961 S3. Doctors and other prescribers should prescribe oxygen using a target saturation range (sections
962 8, 9 and 11) and sign the drug chart or electronic prescribing order. [Grade D]
963

964 S4. An oxygen target saturation range should be prescribed for all patients who are admitted to
965 hospital. This will ensure that every patient will receive appropriate oxygen therapy if it should be
966 required. It will also ensure that all clinicians are aware of the appropriate oxygen target range for
967 every patient under their care. [Grade D]
968

969 **Good practice points related to prescribing and administering oxygen therapy to patients**

- 970 • Oxygen should be prescribed on the drug chart or electronic prescribing system using a
971 target saturation range.
- 972 • In most emergency situations, oxygen is given to patients immediately without a formal
973 prescription. The lack of a prescription should never preclude oxygen being given when
974 needed in an emergency situation. However, a subsequent written record must be made of
975 what oxygen therapy has been given to every patient in a similar manner to the recording of
976 all other emergency treatment.
977

978 **T Monitoring and adjusting therapy (see online sections 9-11)**

979 T1. Pulse oximetry must be available in all locations where emergency oxygen is being used by
980 health care professionals (see also the limitations of using pulse oximetry section 7.1.2). [Grade D]
981

982 T2. All documents which record oximetry measurements or blood gas results should state whether
983 the patient is breathing air or a specified oxygen delivery device and flow rate. [Grade D]
984

985 T3. In all situations where repeated blood gas measurements are required, they should be
986 measured as soon as possible, usually within 30 min of any treatment change, to determine if the
987 proposed target saturations are appropriate. Consider the use of an indwelling arterial catheter if
988 multiple samples are likely to be required. [Grade D]
989

990 T4. Adjustments should only be made by staff who have been trained to administer oxygen. If the
991 oxygen saturation falls below the pre-specified range, the concentration of oxygen should be
992 increased; if the saturation rises above this range, the oxygen concentration should be reduced. If
993 the monitoring of oxygen saturation is performed by other staff (e.g., health care assistants), they
994 should inform staff who are trained to administer oxygen if the oxygen saturation is above or below
995 the target saturation. [Grade D]
996

997 **Good practice points related to administration of oxygen therapy**

- 998 • For hypoxaemic patients, oxygen therapy should continue during other treatments such as
999 nebulised therapy. Clinicians should assess the clinical status of the patient prior to
1000 prescribing oxygen and the patient's condition should be reassessed frequently during
1001 oxygen use (see recommendations B1-B3).
- 1002 • The administering healthcare professional should note the oxygen saturation before
1003 commencing oxygen therapy (see recommendation B3).
- 1004 • The healthcare professional should commence oxygen therapy using an appropriate
1005 delivery system and flow rate as specified in sections 8, 9 and 10 of this guideline. The
1006 target oxygen saturation should be documented on the respiratory section of the
1007 observation chart.

- 1008 • Whenever possible, patients should be given an oxygen information sheet (example in web
1009 appendix 6 of this guideline on the British Thoracic Society Website).

1010
1011

1012 **U Weaning and discontinuation of oxygen therapy**

1013 U1. Lower the oxygen concentration if the patient is clinically stable and the oxygen saturation has
1014 been in the upper zone of the target range for some time (usually 4–8 h). [Grade D]

1015

1016 U2. If the target saturation is maintained, the new delivery system and flow should be continued.
1017 Repeat blood gas measurements are not required. If the patient is stable the process can be
1018 repeated and the patient can eventually be weaned off oxygen (see section 12). [Grade D]

1019

1020 U3. Most stable convalescent patients will eventually be stepped down to 2 l/min via nasal cannulae
1021 prior to cessation of oxygen therapy. Patients at risk of hypercapnic respiratory failure may be
1022 stepped down to 1 l/min (or occasionally 0.5 l/min) via nasal cannulae or a 24% Venturi mask at 2
1023 l/min as the lowest oxygen concentration prior to cessation of oxygen therapy. [Grade D]

1024

1025 U4. Oxygen therapy should be stopped once a patient is clinically stable on low-concentration
1026 oxygen and the oxygen saturation is within the desired range on two consecutive observations (but
1027 the prescription for a target saturation range should remain active in case of future deterioration).
1028 Oxygen should also be stopped if the patient has come to the end of a written protocol of timed
1029 oxygen (e.g. postoperatively). [Grade D]

1030

1031 U5. Oxygen saturation on air should be monitored for 5 min after stopping oxygen therapy. If it
1032 remains in the desired range it should be rechecked at 1 h. [Grade D]

1033

1034 U6. If the oxygen saturation and physiological “track and trigger” score (e.g. NEWS) is satisfactory
1035 at 1 h, the patient has safely discontinued oxygen therapy but saturation and physiology should
1036 continue to be monitored on a regular basis according to the patient’s underlying clinical condition.
1037 [Grade D]

1038

1039 U7. If the saturation falls below the patient’s target range on stopping oxygen therapy,
1040 recommence the lowest concentration that maintained the patient in the target range and monitor
1041 for 5 min. If this restores the saturation into the target range, continue oxygen therapy at this level
1042 and attempt discontinuation of oxygen therapy again at a later date provided the patient remains
1043 clinically stable. [Grade D]

1044

1045 U8. If a patient requires oxygen therapy to be restarted at a higher concentration than before to
1046 maintain the same target saturation range, the patient should have a clinical review to establish the
1047 cause for this deterioration. [Grade D]

1048

1049 U9. Some patients may have episodic hypoxaemia (e.g., after minor exertion or due to mucus
1050 plugging) after they have safely discontinued oxygen therapy. An on-going prescription for a target
1051 saturation range will allow these patients to receive oxygen as the need arises but transient
1052 asymptomatic desaturation does not require correction. [Grade D]

1053

1054

1055

1056

1057

1058 **V Practical aspects of oxygen use in pre-hospital and hospital care and use of oxygen alert**
1059 **cards (see online sections 9-11)**
1060

1061 V1. Emergency oxygen should be available in primary care medical centres, preferably using oxygen
1062 cylinders with integral high-flow regulators. Alternatively, oxygen cylinders fitted with high-flow
1063 regulators (delivering over 6 l/min) must be used. [Grade D]
1064

1065 V2. Trusts should take measures to eliminate the risk of oxygen tubing being connected to the
1066 incorrect wall oxygen outlet or to outlets that deliver compressed air or other gases instead of
1067 oxygen. Air flow meters should be removed from the wall sockets or covered with a designated air
1068 outlet cover when not in use. Special care should be taken if twin oxygen outlets are in use. [Grade
1069 D]
1070

1071 **Good Practice Points related to practical aspects of oxygen therapy**
1072
1073

1074 **Assessment and immediate oxygen therapy**

- 1075 • Chronically hypoxaemic patients with a clinical exacerbation associated with a 3% or greater
1076 fall in oxygen saturation on their usual oxygen therapy should be assessed in hospital with
1077 blood gas estimations. Arterial PO₂ of <7 kPa equates to SpO₂ below approximately 85%.
- 1078 • The initial oxygen therapy to be used in the various clinical situations is given in tables 1–4.
- 1079 • If there is a clear history of asthma or heart failure or other treatable illness, appropriate
1080 treatment should be instituted in accordance with guidelines or standard management plans
1081 for each disease.
- 1082 • The oxygen saturation should be monitored continuously until the patient is stable or
1083 arrives at hospital for a full assessment. The oxygen concentration should be adjusted
1084 upwards or downwards to maintain the target saturation range.
- 1085 • In most emergency situations oxygen is given to patients immediately without a formal
1086 prescription or drug order. The lack of a prescription should never preclude oxygen being
1087 given when needed in an emergency situation. However, a subsequent written record must
1088 be made of what oxygen therapy has been given to every patient (in a similar manner to the
1089 recording of all other emergency treatment).
- 1090 • General practitioners or first responders visiting a patient's home should carry a portable
1091 pulse oximeter to assess hypoxaemia and guide use of oxygen if available.
- 1092 • Those attending patients as an emergency in rural or remote areas should consider carrying
1093 a portable oxygen cylinder and appropriate delivery systems as part of their emergency
1094 equipment.

1095
1096
1097 **Oxygen alert cards for patients with hypercapnic respiratory failure**

- 1098 • Patients with COPD (and other at-risk conditions) who have had an episode of hypercapnic
1099 respiratory failure should be issued with an oxygen alert card and with a 24% or 28% Venturi
1100 mask. They should be instructed to show the card to the ambulance crew and emergency
1101 department staff in the event of an exacerbation.
- 1102 • Oxygen alert cards with agreed content can be obtained via the British Thoracic Society (see
1103 Emergency Oxygen Guideline Section on BTS Website)
- 1104 • The content of the alert card should be specified by the physician in charge of the patient's
1105 care, based on previous blood gas results.
- 1106 • The primary care team and ambulance service should also be informed by the hospital COPD
1107 team that the patient has had an episode of hypercapnic respiratory failure and carries an
1108 oxygen alert card. The home address and ideal oxygen concentration or target saturation

1109 ranges of these patients can be flagged in the ambulance control systems and information
1110 disseminated to ambulance crews when required.
1111 • When possible, out-of-hours services providing emergency primary care services should be
1112 informed by the hospital COPD team or by the primary care team that the patient has had
1113 an episode of hypercapnic respiratory failure and carries an oxygen alert card. Use of oxygen
1114 in these patients will be guided by the instructions on the alert card.

1115

1116 **W Practical aspects of oxygen dispensing, documentation and monitoring**

1117

1118 W1. Nurses and others who dispense drugs in hospitals should sign the drug chart or electronic
1119 prescribing record at every drug round and check that the patient is receiving oxygen therapy. This is
1120 to check that the patient is within the target saturation and also to check whether weaning and
1121 discontinuation should be instituted. [Grade D]

1122

1123 W2. Most patients are prescribed an oxygen target range. If patients are on air at the time of the
1124 drug round, nurses should sign the drug chart using a code such as “A” for air and the observation
1125 chart should also be filled in using the code A for air (see chart 4, fig 4). [Grade D]

1126

1127 W3. All patients should have their oxygen saturation observed for at least five minutes after starting
1128 oxygen therapy or for patients who require an increased concentration of oxygen and after oxygen
1129 therapy has been decreased or stopped. [Grade D]

1130

1131 W4. If the oxygen saturation is above the target saturation range and the patient is stable, the
1132 delivery system or oxygen flow rate should be modified to return the saturation to within the target
1133 range. [Grade D]

1134

1135 W5. Patients who have a target saturation of 88–92% should have their blood gases measured
1136 within 30–60 min. This is to ensure that the carbon dioxide level is not rising. This recommendation
1137 also applies to those who are at risk of developing hypercapnic respiratory failure but who have a
1138 normal PaCO₂ on the initial blood gas measurement. [Grade D]

1139

1140 W6. Stable patients whose oxygen saturation is within their target saturation range of 94–98% do
1141 not need repeat blood gas measurements within 30–60 min if there is no risk of hypercapnic
1142 respiratory failure and acidosis and may not need any further blood gas measurements. [Grade D]

1143

1144 W7. Stable patients on oxygen treatment should have SpO₂ and physiological variables (e.g., NEWS)
1145 measured four times a day. [Grade D]

1146

1147 W8. In those who are unstable (e.g. NEWS score 7 or above), oxygen saturation should be
1148 monitored continuously and the patient may require level 2 or level 3 care on a high dependency
1149 unit or critical care unit. [Grade D]

1150

1151 W9. If the patient is clinically stable and the oxygen saturation is within the target range, treatment
1152 should be continued (or eventually lowered) depending on the clinical situation. [Grade D]

1153

1154 W10. Oxygen therapy should be increased if the saturation is below the desired range and
1155 decreased if the saturation is above the desired range (and eventually discontinued as the patient
1156 recovers). [Grade D]

1157

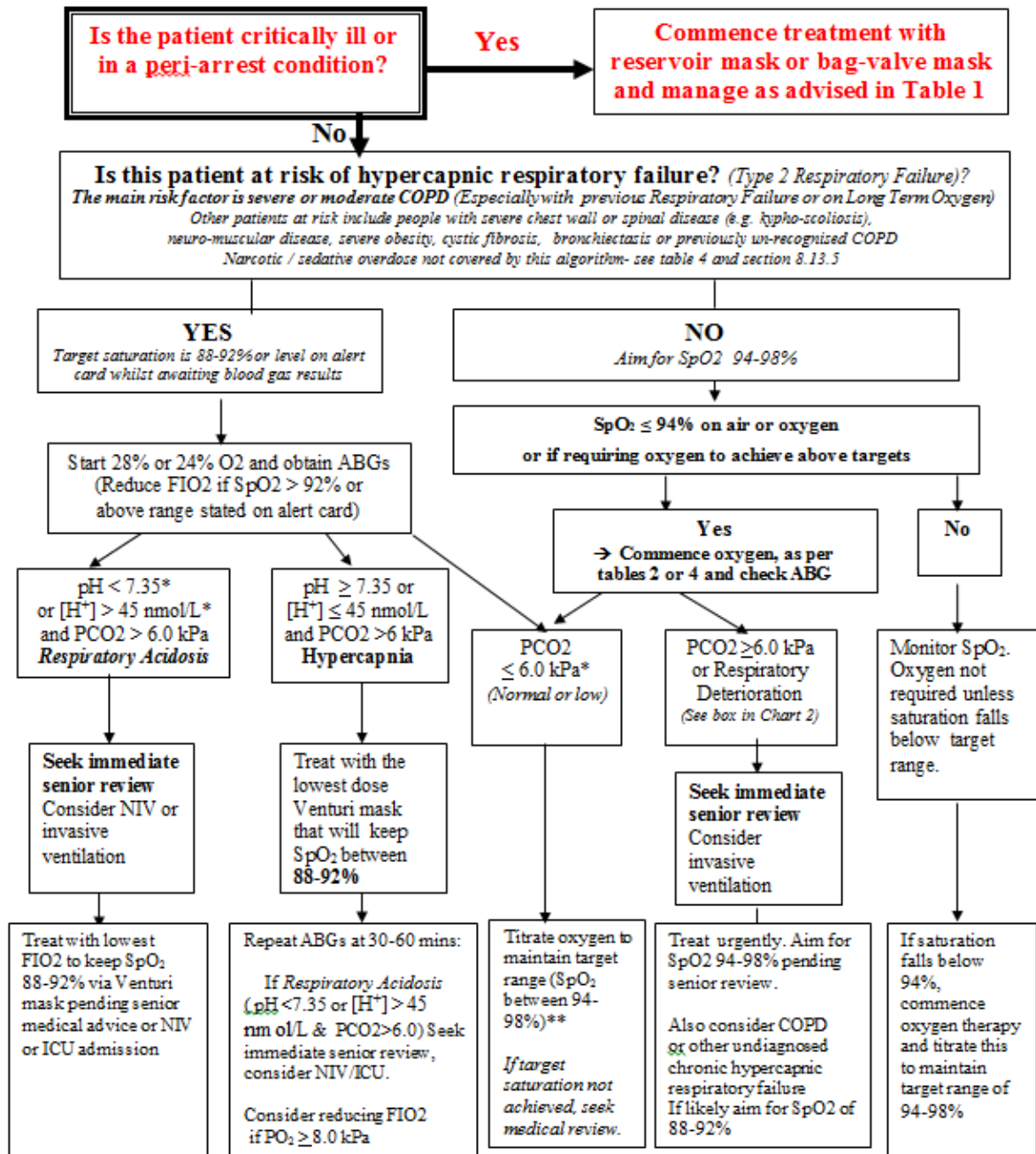
1158 W11. The new saturation (and the new delivery system) and flow rate should be recorded on the
1159 patient’s observation chart after 5 min of treatment at the new oxygen concentration. Each change

1160 should be recorded by the clinician trained to administer oxygen by signing the observation chart
1161 (only changes should be signed for). [Grade D]
1162
1163 W12. Repeat blood gas measurements are not required for stable patients who require a reduced
1164 concentration of oxygen (or cessation of oxygen therapy) to maintain the desired target saturation.
1165 [Grade D]
1166
1167 W13. Patients with no risk of hypercapnic respiratory failure do not always need repeat blood gas
1168 measurements after an increase in oxygen concentration. However, the patient requires clinical
1169 review to determine why the oxygen saturation has fallen. [Grade D]
1170
1171 W14. Patients at risk of hypercapnic respiratory failure (usually those with a target range of 88–
1172 92%; see table 4) require repeat blood gas estimation 30–60 min after an increase in oxygen therapy
1173 (to ensure that the carbon dioxide level is not rising). [Grade D]
1174
1175 W15. For patients with no risk of hypercapnic respiratory failure, monitoring by pulse oximeter is
1176 sufficient (repeated blood gases not required) provided the saturation remains in the desired range,
1177 usually 94–98%. [Grade D]
1178
1179 W16. If a patient’s oxygen saturation is lower than the prescribed target range, first check all
1180 aspects of the oxygen delivery system and the oximeter device for faults or errors. [Grade D]
1181
1182 W17. If a patient’s oxygen saturation is consistently lower than the prescribed target range, there
1183 should be a medical review and the oxygen therapy should be increased according to an agreed
1184 written protocol. [Grade D]
1185
1186 W18. If the oxygen saturation fails to rise following 5– 10 min of increased oxygen therapy or if
1187 there is clinical concern following medical review, then blood gas measurements should be
1188 repeated. [Grade D]
1189
1190

X Training in oxygen prescribing and use

1191
1192 X1. All clinicians prescribing oxygen should have appropriate training and access to written or
1193 electronic oxygen prescribing guidelines based on this national guideline. [Grade D]
1194
1195 X2. Every hospital should have a training programme to ensure that clinical staff are familiar with
1196 the hospital’s oxygen administration policies. In view of the high number of adverse incidents
1197 related to oxygen therapy, it is recommended that all acute Trusts should include basic training in
1198 oxygen use in the mandatory training programmes for all clinical staff. [Grade D]
1199

Chart 1 Oxygen prescription for acutely hypoxaemic patients in hospital



Any increase in FIO2 must be followed by repeat ABGs in 1 hour (or sooner if conscious level deteriorates)
 * If pH is < 7.35 ([H+] > 45 nmol/L) with normal or low PaCO2, investigate and treat for metabolic acidosis and keep SpO2 94-98%
 **Repeat ABC in 30-60 mins for all patients at risk of Type 2 respiratory failure (even if initial PaCO2 is normal)

1200
1201
1202
1203
1204

Chart 2

Flow chart for oxygen administration on general wards in hospitals

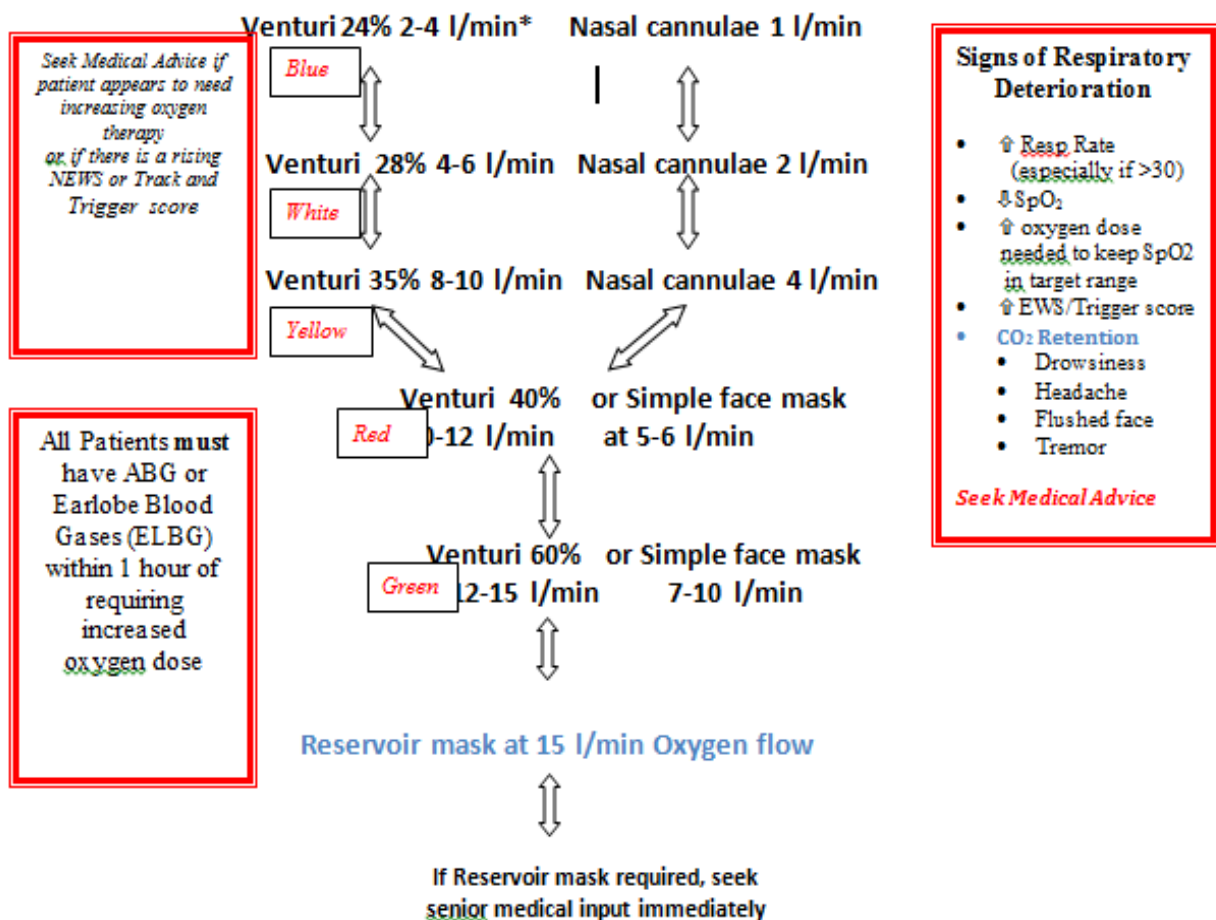
See patient's drug chart and Chart 1 and tables 1-4 for starting dose and target saturation
Choose the most suitable delivery system and flow rate

Titrate oxygen up or down to maintain the target oxygen saturation.

The table below shows available options for stepping dosage up or down. The chart does NOT imply any equivalence of dose between Venturi masks and nasal cannulae.

Allow at least 5 minutes at each dose before adjusting further upwards or downwards (except with major and sudden fall in saturation)

Once your patient has adequate and stable saturation on minimal oxygen dose, consider discontinuation of oxygen therapy.



* For Venturi masks, the higher flow rate is required if the respiratory rate is >30

Patients in a peri-arrest situation and critically ill patients should be given maximal oxygen therapy via reservoir mask or bag-valve mask whilst immediate medical help is arriving.

(Except for patients with COPD with known oxygen sensitivity recorded in patient's case notes and drug chart or in the EPR: keep saturation at 88-92% for this sub-group of patients)

1206 **Table 5: SIGN evidence levels**

SIGN LEVELS OF EVIDENCE

1++	High quality meta-analyses, systematic reviews of RCTs, or RCTs with a very low risk of bias
1+	Well-conducted meta-analyses, systematic reviews, or RCTs with a low risk of bias
1-	Meta-analyses, systematic reviews, or RCTs with a high risk of bias
2++	High quality systematic reviews of case control or cohort studies High quality case control or cohort studies with a very low risk of confounding or bias and a high probability that the relationship is causal
2+	Well-conducted case control or cohort studies with a low risk of confounding or bias and a moderate probability that the relationship is causal
2-	Case control or cohort studies with a high risk of confounding or bias and a significant risk that the relationship is not causal
3	Non-analytic studies, e.g. case reports, case series
4	Expert opinion

1207

1208

1209 **Table 6: SIGN grades of recommendation**

1210

GRADES OF RECOMMENDATIONS

- A** At least one meta-analysis, systematic review, or RCT rated as 1++, and directly applicable to the target population; or
A body of evidence consisting principally of studies rated as 1+, directly applicable to the target population, and demonstrating overall consistency of results
- B** A body of evidence including studies rated as 2++, directly applicable to the target population, and demonstrating overall consistency of results; or
Extrapolated evidence from studies rated as 1++ or 1+
- C** A body of evidence including studies rated as 2+, directly applicable to the target population and demonstrating overall consistency of results; or
Extrapolated evidence from studies rated as 2++
- D** Evidence level 3 or 4; or
Extrapolated evidence from studies rated as 2+

Good practice points

- Recommended best practice based on the clinical experience of the guideline development group

1211

1212

Public consultation



1213
1214
1215
1216
1217
1218
1219
1220
1221
1222
1223
1224
1225
1226
1227
1228
1229
1230
1231
1232
1233
1234
1235
1236
1237
1238
1239
1240
1241

References

1. O'Driscoll BR, Howard LS, Davison AG, et al. BTS guideline for emergency oxygen use in adult patients. *Thorax* 2008;**63 Suppl 6**:vi1-68 doi: 10.1136/thx.2008.102947[published Online First: Epub Date]].
2. Hardinge M, Annandale J, Bourne S, et al. British Thoracic Society guidelines for home oxygen use in adults. *Thorax* 2015;**70 Suppl 1**:i1-43 doi: 10.1136/thoraxjnl-2015-206865[published Online First: Epub Date]].
3. Hale KE, Gavin C, O'Driscoll BR. Audit of oxygen use in emergency ambulances and in a hospital emergency department. *Emerg Med J* 2008;**25**(11):773-6 doi: 10.1136/emj.2008.059287[published Online First: Epub Date]].
4. Wilson AT, Channer KS. Hypoxaemia and supplemental oxygen therapy in the first 24 hours after myocardial infarction: the role of pulse oximetry. *J R Coll Physicians Lond* 1997;**31**(6):657-61
5. Cook DJ, Reeve BK, Griffith LE, et al. Multidisciplinary education for oxygen prescription. A continuous quality improvement study. *Arch Intern Med* 1996;**156**(16):1797-801
6. Small D, Duha A, Wieskopf B, et al. Uses and misuses of oxygen in hospitalized patients. *Am J Med* 1992;**92**(6):591-5
7. Dodd ME, Kellet F, Davis A, et al. Audit of oxygen prescribing before and after the introduction of a prescription chart. *BMJ* 2000;**321**(7265):864-5
8. Howell M. An audit of oxygen prescribing in acute general medical wards. *Prof Nurse* 2001;**17**(4):221-4
9. Boyle M, Wong J. Prescribing oxygen therapy. An audit of oxygen prescribing practices on medical wards at North Shore Hospital, Auckland, New Zealand. *N Z Med J* 2006;**119**(1238):U2080
10. Kbar FA, Campbell IA. Oxygen therapy in hospitalized patients: the impact of local guidelines. *J Eval Clin Pract* 2006;**12**(1):31-6 doi: 10.1111/j.1365-2753.2005.00601.x[published Online First: Epub Date]].
11. AGREE II Instrument. Secondary AGREE II Instrument. <http://www.agreetrust.org/agree-ii/>.
12. JRCALC. *UK Ambulance Services Clinical Practice Guidelines 2013*. London: JRCALC, 2013.
13. BTS. *British Thoracic Society Guideline Production Manual*. London: BTS, 2014:31.