

National COPD Audit Programme



COPD: Working together

National Chronic Obstructive Pulmonary Disease (COPD) Audit Programme: Clinical audit of COPD exacerbations admitted to acute hospitals in England and Wales 2017

Data analysis and methodology
April 2018

Prepared by:



**Royal College
of Physicians**

**Imperial College
London**

In partnership with:



**British
Thoracic
Society**

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Working in wider partnership with:



The Royal College of Physicians

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Healthcare Quality Improvement Partnership (HQIP)

The National COPD Audit Programme is commissioned by the Healthcare Quality Improvement Partnership (HQIP) as part of the National Clinical Audit (NCA) Programme. HQIP is led by a consortium of the Academy of Medical Royal Colleges, the Royal College of Nursing and National Voices. Its aim is to promote quality improvement, and in particular to increase the impact that clinical audit has on healthcare quality in England and Wales. HQIP holds the contract to manage and develop the NCA Programme, comprising more than 30 clinical audits that cover care provided to people with a wide range of medical, surgical and mental health conditions. The programme is funded by NHS England, the Welsh Government and, with some individual audits, also funded by the Health Department of the Scottish Government, DHSSPS Northern Ireland and the Channel Islands.

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How to use this report

This report presents the results from an extensive analysis of the data from the clinical component of the National Chronic Obstructive Pulmonary Disease (COPD) Audit Programme's secondary care audit. Details of the methodology employed are also provided ([Appendix C](#)).

The continuous audit, which captures the process and clinical outcomes of treatment in patients admitted to hospital in England and Wales with COPD exacerbations, launched on 1 February 2017. This report, which is the first report post launch of continuous data collection, presents the results of the cohort of patients discharged between the audit's launch date and 13 September 2017. In order for the report to be published in line with the National COPD Audit Programme's contract end date, the data were extracted prior to completion of a full year of data collection. Consequently, this report presents the results of a slightly truncated patient cohort.

References to the appropriate National Institute for Health and Care Excellence (NICE) quality statements^{a,b} ([Appendix D](#)) and clinical guidelines^c ([Appendix E](#)) are inserted throughout the key findings.

The data are presented largely in tabular form with explanatory notes where appropriate. Although these data are available to the interested reader, it is not necessary to review them to appreciate the key messages, which can be found in the national report (via www.rcplondon.ac.uk/working-together). Direct comparisons to 2014 audit data have been provided, where appropriate. Where in some instances data is not directly comparable, 2014 data have been presented in a separate table beneath the 2017 audit data.

Copies of our datasets, our good practice repository as well as the resources supplied for both the clinical and organisational audits can be found via our website: www.rcplondon.ac.uk/projects/outputs/secondary-care-audit-2017-resources.

The Best practice tariff (BPT) for COPD is in place for the financial years 2017/18 and 2018/19 for trusts in England. Publically available BPT quarterly and year to date reports can be downloaded online from: www.nacap.org.uk/nacap/welcome.nsf/reportsSC.html. In the future regional reports will also be made available on a six monthly basis.

If you would like to discuss any of the findings or recommendations in more detail with other participants, or you have ideas you would like to share, you can log into our audit forum on the Respiratory Futures website: www.respiratoryfutures.org.uk/copdsecondarycareauditforum/.

^a National Institute for Health and Care Excellence. *Chronic obstructive pulmonary disease in adults*. NICE Quality standard 10 (QS10). London: NICE, 2016. www.nice.org.uk/Guidance/QS10 [Accessed December 2017].

^b National Institute for Health and Care Excellence. *Chronic obstructive pulmonary disease in adults*. NICE Quality standard 10 (QS10). 2011 edition. London: NICE, 2011. www.nice.org.uk/Guidance/QS10 [Accessed December 2017].

^c National Institute for Health and Care Excellence. *Chronic obstructive pulmonary disease in over 16s: diagnosis and management* (partial update). NICE Clinical Guideline 101 (CG101). London: NICE, 2010. www.nice.org.uk/guidance/CG101 [Accessed December 2017].



Section 1: General information

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Key findings

Admissions/discharge

- The median **time from arrival to admission** was **3.4 hours**.^d
 - Admission rates** for COPD were **greater during weekdays** than at weekends. The **highest percentage** of admissions were on a **Monday (15.7%)**.^e
- As per the results of the 2014 audit, **fewer patients** were **discharged on a weekend** (8.4% on a Saturday and 6.4% on a Sunday in 2017, 7% on a Saturday and 5% on a Sunday in 2014), in comparison to during the weekday (for example, 15.9% on a Monday and 17.8% on a Tuesday in 2017, 16% on a Monday and 19% on a Tuesday in 2014).

Length of stay

- The **median length of stay** remained unchanged from that reported in 2014, at **4 days**.

Mortality

- Inpatient mortality fell** marginally (3.9% reported in 2017 vs 4.3% reported in 2014).

For the full key findings and recommendations, please see the national report *COPD: Working together*, available at www.rcplondon.ac.uk/working-together

Navigation

This section contains the following tables and graphs. If you are viewing this report on a computer, you can select the table that you wish to see from the list below.

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 - [1.4.1 Average number of admissions per hospital](#)
 - [1.4.2 Age at admission by gender](#)
 - [1.4.3 Average time, in hours, between arrival and admission](#)
 - [1.4.4 Day and time of admission to hospital](#)

1.1 Age

| Age at admission | 2017 (N=36,341) | 2014 (N=13,414) |
|---------------------|-----------------|-----------------|
| Mean | 71.9 | 72 |
| Standard deviation | 10.7 | 11 |
| Median | 73.0 | 72 |
| Interquartile range | 65–80 | 65–80 |

^d There is no comparative data from 2014, as only date and time of admission were captured in that audit (ie not of arrival).

^e Comparative data from 2014 cannot be reported here as admission was calculated from time recorded of arrival to the unit.

1.2 Gender

| Gender | 2017 (N=36,341) | 2014 (N=13,414) |
|--------|-----------------|-----------------|
| Male | 46.9% (17,046) | 49% (6,572) |
| Female | 53.1% (19,295) | 51% (6,842) |

1.3 Socioeconomic status

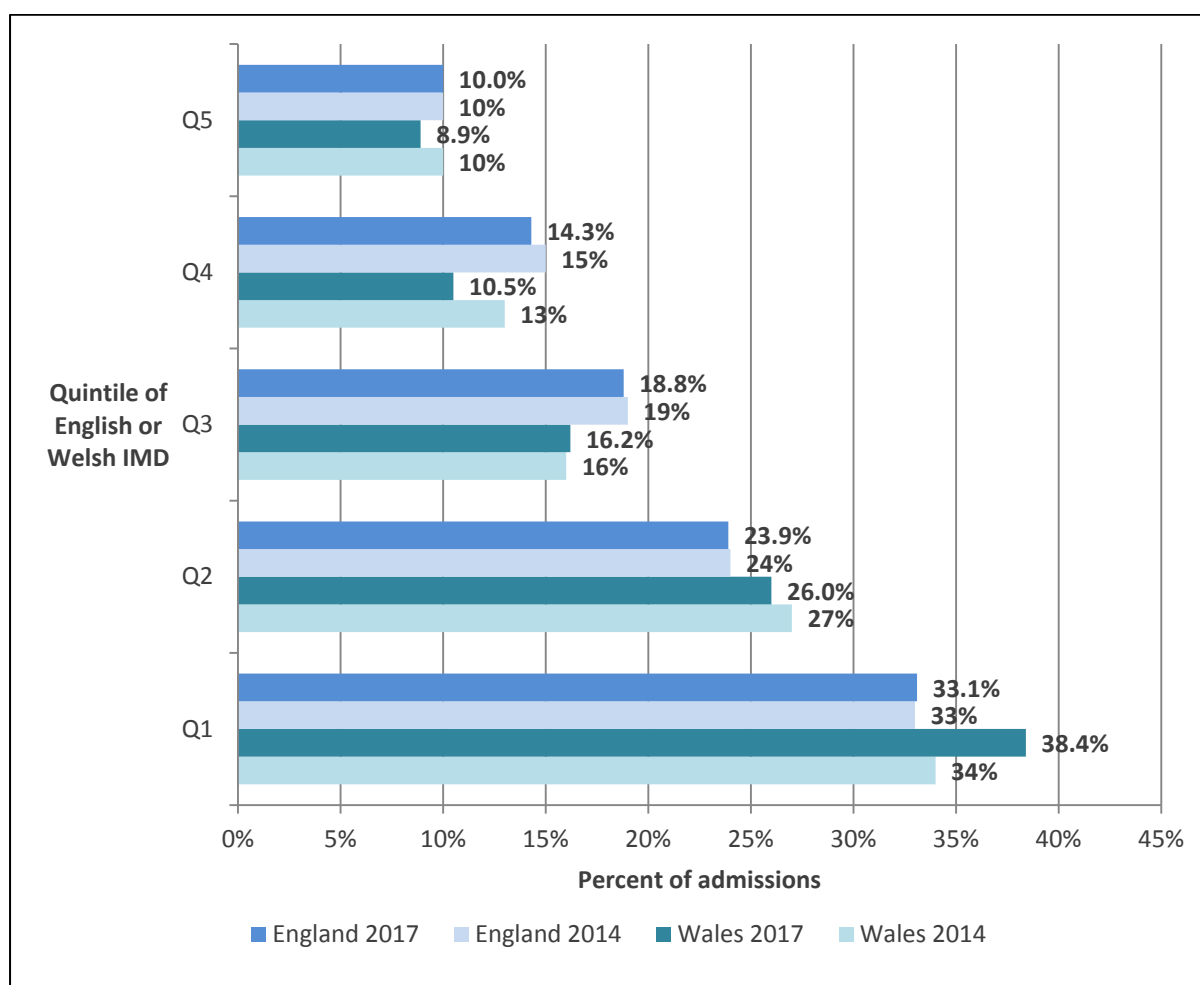
1.3.1 Index of Multiple Deprivation measures by national quintile in England

| Index of Multiple Deprivation | % of audit sample living in an English Lower-layer Super Output Area (LSOA) | | | | |
|-------------------------------|---|---------------|---------------|---------------|---------------------|
| | Q1 (Most deprived) | Q2 | Q3 | Q4 | Q5 (Least deprived) |
| England 2017 (N=34,327) | 33.1% (11,363) | 23.9% (8,196) | 18.8% (6,452) | 14.3% (4,902) | 10.0% (3,414) |
| England 2014 (N=12,245) | 33% (4,006) | 24% (2,895) | 19% (2,282) | 15% (1,804) | 10% (1,258) |

1.3.2 Index of Multiple Deprivation measures by national quintile in Wales

| Index of Multiple Deprivation | % of audit sample living in a Welsh Lower-layer Super Output Area (LSOA) | | | | |
|-------------------------------|--|-------------|-------------|-------------|---------------------|
| | Q1 (Most deprived) | Q2 | Q3 | Q4 | Q5 (Least deprived) |
| Wales 2017 (N=1,664) | 38.4% (639) | 26.0% (433) | 16.2% (269) | 10.5% (175) | 8.9% (148) |
| Wales 2014 (N=829) | 34% (283) | 27% (224) | 16% (135) | 13% (108) | 10% (79) |

Deprivation quintile of admissions to hospital with a COPD exacerbation in England and Wales

1.4 Admission^f

1.4.1 Average number of admissions per hospital

| Number of admissions | 2017 (N=36,341) | 2014 (N=13,414) |
|----------------------|-----------------|-----------------|
| Mean | 201.8 | Not reported |
| Standard deviation | 144.1 | Not reported |
| Median | 179 | 61 |
| Interquartile range | 94–263 | 38–85 |

1.4.2 Age at admission by gender

| Age at admission by gender | 2017 (N=36,341) | | 2014 (N=13,414) | |
|----------------------------|-----------------|-------------------|-----------------|------------------|
| | Male (N=17,046) | Female (N=19,295) | Male (N=6,572) | Female (N=6,842) |
| Mean | 72.1 | 71.8 | 72 | 72 |
| Standard deviation | 10.6 | 10.9 | Not reported | Not reported |

^f Question 1.6 from the dataset was: Date of and time of [patient] arrival at your hospital, and question 1.7 was: Date and time of [patient] admission to the unit. Therefore 1.4 here combines questions 1.6 and 1.7 from the dataset.

1.4.3 Average time, in hours, between arrival and admission

| Time of arrival to admission, in hours | 2017 (N=36,341) |
|--|-----------------|
| Mean | 3.9 |
| Standard deviation | 3.6 |
| Median | 3.4 |
| Interquartile range | 1.2–5.1 |

1.4.4 Day and time of admission to hospital

| Time admitted | Day patient admitted (N=36,341) | | | | | | |
|-----------------|---------------------------------|----------------------|------------------------|-----------------------|---------------------|-----------------------|---------------------|
| | Monday (N=5,710) | Tuesday (N=5,463) | Wednesday (N=5,472) | Thursday (N=5,385) | Friday (N=5,260) | Saturday (N=4,433) | Sunday (N=4,618) |
| 00.00– 01.59 | 7.2% (409) | 8.2% (447) | 8.1% (442) | 7.4% (396) | 8.4% (442) | 9.0% (397) | 8.6% (396) |
| 02.00– 03.59 | 5.5% (312) | 6.8% (373) | 6.5% (356) | 6.3% (341) | 6.2% (325) | 8.2% (363) | 6.9% (320) |
| 04.00– 05.59 | 4.4% (250) | 5.0% (274) | 4.4% (241) | 4.6% (250) | 4.5% (238) | 6.3% (277) | 5.4% (250) |
| 06.00– 07.59 | 3.7% (210) | 4.0% (219) | 3.9% (214) | 4.0% (213) | 3.9% (206) | 4.0% (177) | 4.3% (198) |
| 08.00– 09.59 | 5.4% (308) | 5.3% (289) | 5.5% (300) | 4.9% (266) | 4.9% (255) | 5.9% (262) | 5.3% (243) |
| 10.00– 11.59 | 7.4% (423) | 8.2% (447) | 7.3% (398) | 7.7% (414) | 7.8% (410) | 7.9% (349) | 7.7% (355) |
| 12.00– 13.59 | 9.9% (566) | 9.8% (534) | 9.3% (509) | 9.2% (495) | 9.3% (490) | 8.7% (386) | 9.2% (423) |
| 14.00– 15.59 | 12.4% (710) | 11.8% (644) | 11.6% (637) | 12.1% (652) | 11.7% (617) | 10.7% (474) | 10.9% (504) |
| 16.00– 17.59 | 12.5% (716) | 11.9% (649) | 11.8% (647) | 13.3% (717) | 11.5% (604) | 10.2% (452) | 12.1% (557) |
| 18.00– 19.59 | 12.1% (692) | 11.1% (608) | 11.5% (629) | 11.5% (619) | 10.8% (566) | 10.0% (442) | 9.7% (450) |
| 20.00– 21.59 | 9.4% (537) | 9.5% (521) | 9.9% (539) | 9.3% (502) | 10.1% (532) | 9.7% (431) | 9.7% (450) |
| 22.00– 23.59 | 10.1% (577) | 8.4% (458) | 10.2% (560) | 9.7% (520) | 10.9% (575) | 9.5% (423) | 10.2% (472) |



Section 2: Provision of timely care

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Key findings

Acute physician review ([NICE \[CG101\] 1.1.8.2](#))

- **82.3% of admissions** were reviewed by an acute physician of grade specialty trainee 3 (ST3) or above.^g

Respiratory specialist review ([NICE \[QS10\], statement 10, 2011](#))

- **78% of admissions** were reviewed by a member of the **respiratory team**.
- The median time from admission to review by a member of the specialist respiratory team has reduced (16.2 hours in 2017, compared to 21 hours in 2014^h).
- There was an improvement in the number of admissions **reviewed by a member of the respiratory team within 24 hours** (54.8% in 2017, compared to 49% in 2014).

For the full key findings and recommendations, please see the national report *COPD: Working together*, available at www.rcplondon.ac.uk/working-together

Navigation

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2.1 Acute physician review

2.1.1 Has the patient been reviewed by an acute physician of grade ST3 or above?

| Review by an acute physician of grade ST3 or above | 2017 (N=36,341) |
|--|-----------------|
| No | 12.8% (4,635) |
| Yes | 82.3% (29,919) |
| Not recorded | 4.9% (1,787) |

^g This was a new question in the 2017 dataset and, therefore, there is no comparative data from the 2014 audit.

^h Seen by a respiratory consultant or a respiratory nurse/member of the COPD/respiratory team.

2.2 Respiratory team review

2.2.1 Has a member of the respiratory team reviewed the patient during the admission?

| Respiratory team review during admission | 2017 (N=36,341) | 2014 (N=13,414) |
|--|-----------------|---------------------------|
| No | 22.0% (7,981) | 20% (2,688) |
| Yes | 78.0% (28,360) | 77% (10,387) ⁱ |
| Not recorded | Not asked | 3% (339) |

2.2.2 Average time, in hours, from admission to respiratory team review

| Time, in hours, from admission to respiratory team review | 2017 (N=36,341) | 2014 (N=8,884) ^j |
|---|-----------------|-----------------------------|
| Mean | 27.0 | Not reported |
| Standard deviation | 50.3 | Not reported |
| Median | 16.2 | 21 |
| Interquartile range | 7.0–30.4 | 12–45 |

2.2.3 Day and time of review by a member of the respiratory team

| Time of review | Day of review (N=28,360) | | | | | | |
|----------------|--------------------------|-------------------|---------------------|--------------------|------------------|--------------------|------------------|
| | Monday (N=5,550) | Tuesday (N=5,059) | Wednesday (N=4,389) | Thursday (N=4,510) | Friday (N=4,450) | Saturday (N=2,339) | Sunday (N=2,063) |
| 00.00–01.59 | 0.7% (39) | 0.6% (31) | 0.6% (28) | 0.9% (41) | 0.6% (26) | 1.2% (27) | 1.7% (34) |
| 02.00–03.59 | 0.4% (21) | 0.6% (32) | 0.6% (24) | 0.4% (19) | 0.5% (23) | 1.2% (28) | 0.8% (16) |
| 04.00–05.59 | 0.4% (21) | 0.4% (21) | 0.6% (28) | 0.4% (20) | 0.4% (17) | 1.0% (24) | 1.3% (26) |
| 06.00–07.59 | 0.7% (37) | 0.9% (44) | 0.6% (25) | 0.8% (35) | 0.7% (32) | 1.3% (30) | 1.8% (37) |
| 08.00–09.59 | 21.7% (1,206) | 22.5% (1,136) | 21.7% (953) | 20.1% (907) | 23.3% (1,035) | 25.4% (594) | 27.1% (560) |
| 10.00–11.59 | 40.8% (2,266) | 38.8% (1,962) | 39.0% (1,712) | 40.2% (1,813) | 37.2% (1,655) | 32.0% (749) | 28.4% (585) |
| 12.00–13.59 | 13.2% (733) | 12.4% (625) | 12.2% (537) | 12.2% (548) | 12.9% (575) | 12.7% (296) | 12.3% (253) |
| 14.00–15.59 | 11.5% (639) | 12.5% (630) | 12.6% (553) | 12.4% (560) | 12.7% (563) | 9.7% (226) | 9.6% (198) |
| 16.00–17.59 | 6.0% (331) | 6.4% (323) | 6.2% (272) | 6.6% (296) | 5.9% (261) | 6.3% (148) | 8.2% (170) |
| 18.00–19.59 | 2.6% (143) | 2.9% (148) | 3.3% (144) | 3.8% (171) | 2.9% (128) | 5.2% (121) | 5.0% (103) |
| 20.00–21.59 | 1.4% (75) | 1.3% (68) | 1.6% (72) | 1.1% (51) | 2.0% (91) | 2.7% (62) | 2.1% (43) |
| 22.00–23.59 | 0.7% (39) | 0.8% (39) | 0.9% (41) | 1.1% (49) | 1.0% (44) | 1.5% (34) | 1.8% (38) |

ⁱ Seen by a respiratory consultant or a respiratory nurse/member of the COPD/respiratory team

^j Seen by a respiratory consultant or a respiratory nurse/member of the COPD/respiratory team

2.2.4 Was the patient reviewed by a member of the respiratory team within 24 hours?

| Respiratory team review within 24 hours of admission | 2017 (N=36,341) | 2014 (N=10,387) |
|---|------------------------|------------------------|
| No | 45.2% (16,414) | 37% (3,789) |
| Yes | 54.8% (19,927) | 49% (5,095) |
| No time seen recorded | Not asked | 15% (1,503) |



Section 3: Recording key clinical information

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Key findings

Oxygen prescription ([NICE \[CG101\] 1.3.6](#)/[NICE \[QS10\] statement 6](#))

- There was a marginal improvement in the number of admissions being prescribed oxygen; **57.3% in 2017**, compared to 55% in 2014.

Spirometry ([NICE \[CG101\] 1.1.2](#)/[NICE \[QS10\] statement 1](#))

- A clear problem was identified with the recording/noting of spirometry. A **spirometry result** was available for only **39.7% of admissions in 2017**, compared to 46% in 2014.
- Importantly, 12.4% of patient admissions in whom spirometry was recorded had no evidence of airflow obstruction ($FEV_1/FVC \geq 0.7$), despite being managed for COPD exacerbation.

Smoking cessation ([NICE \[CG101\] 1.2.1](#))

- The recording of **smoking status** has not improved (**9.1% of admissions** in 2017 were **not asked** about their smoking status/it was not recorded by hospitals, compared to 8% in 2014).
- There was a **reduction** in the number of **self-reported current smokers** (31.3% in 2017, compared to 34% in 2014).

For the full key findings and recommendations, please see the national report *COPD: Working together*, available at www.rcplondon.ac.uk/working-together

Navigation

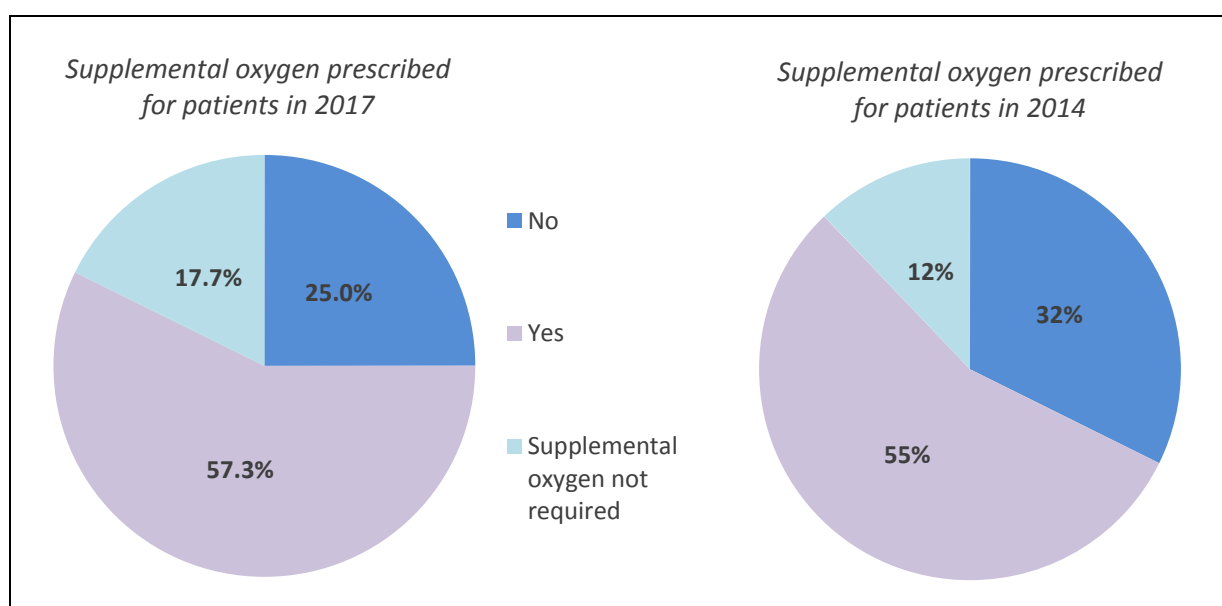
This section contains the following tables and graphs. If you are viewing this report on a computer, you can select the table that you wish to see from the list below.

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 - [3.4.1 Was a DECAF score recorded for this patient?](#)
 - [3.4.2 If yes, what was the recorded DECAF score?](#)

3.1. Oxygen

3.1.1 Was oxygen prescribed for this patient?

| Oxygen prescribed | 2017 (N=36,341) | 2014 (N=13,414) ^k |
|----------------------------------|-----------------|------------------------------|
| No | 25.0% (9,081) | 32% (4,313) |
| Yes | 57.3% (20,829) | 55% (7,434) |
| Supplemental oxygen not required | 17.7% (6,431) | 12% (1,667) |



3.1.2 If oxygen was prescribed was it to a stipulated target range?

| Target range for oxygen prescription | 2017 (N=20,829) | 2014 (N=7,434) |
|--------------------------------------|-----------------|----------------|
| 88–92% | 85.1% (17,733) | 84% (6,251) |
| 94–98% | 8.2% (1,700) | 8% (563) |
| Other | 3.5% (729) | 3% (230) |
| Target range not stipulated | 3.2% (667) | 2% (183) |

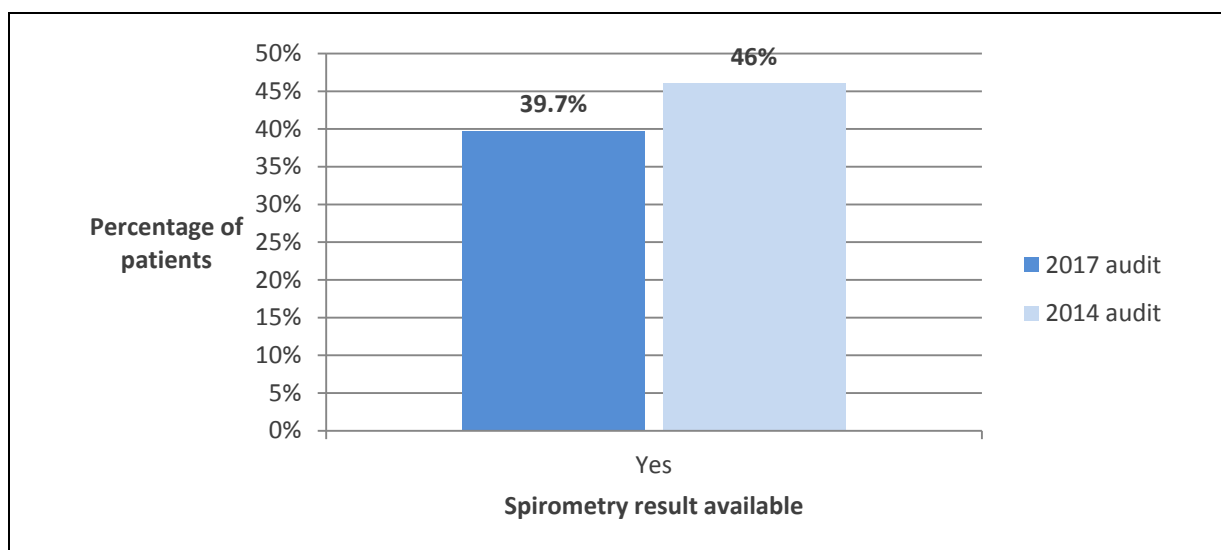
3.2 Spirometric evidence of COPD

3.2.1 Is a spirometry result available?

| Spirometry result available | 2017 (N=36,341) | 2014 (N=13,414) |
|-----------------------------|-----------------|-----------------|
| No | 60.4% (21,931) | 54% (7,291) |
| Yes | 39.7% (14,410) | 46% (6,123) |

^k In 2014 this question was asked as follows: *Was oxygen prescribed on the medication chart or equivalent during this admission?* The options to answer were: *yes, no, and not required*

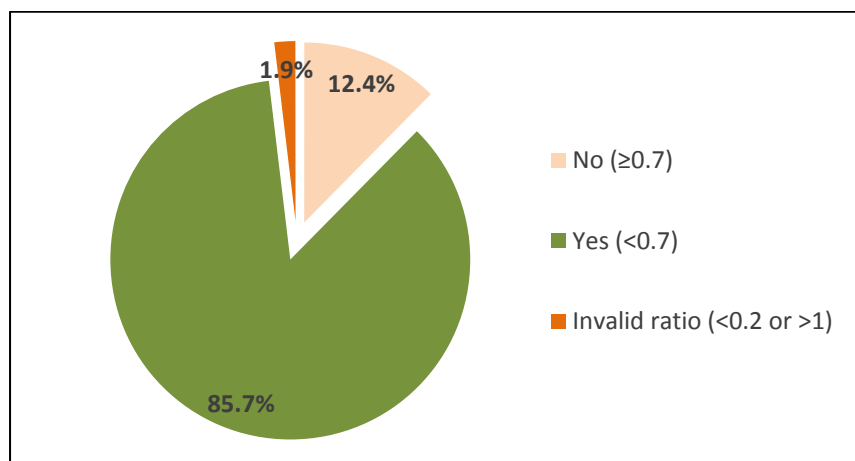
Availability of spirometry results for admissions in the 2017 and 2014 audits



3.2.2 Airflow obstruction¹

| Airflow obstruction (FEV ₁ /FVC ratio) | 2017 (N=14,410) |
|---|-----------------|
| No (≥0.7) | 12.4% (1,783) |
| Yes (<0.7) | 85.7% (12,354) |
| Invalid ratio (<0.2 or >1) | 1.9% (273) |

Evidence of airflow obstruction for those patients with a spirometry result recorded



3.2.3 If a spirometry result is available, what is the patients most recent FEV₁?

| Patient's most recent FEV ₁ , in litres | 2017 (N=14, 410) |
|--|------------------|
| Mean | 1.1 |
| Standard deviation | 0.6 |
| Median | 0.9 |
| Interquartile range | 0.7–1.3 |

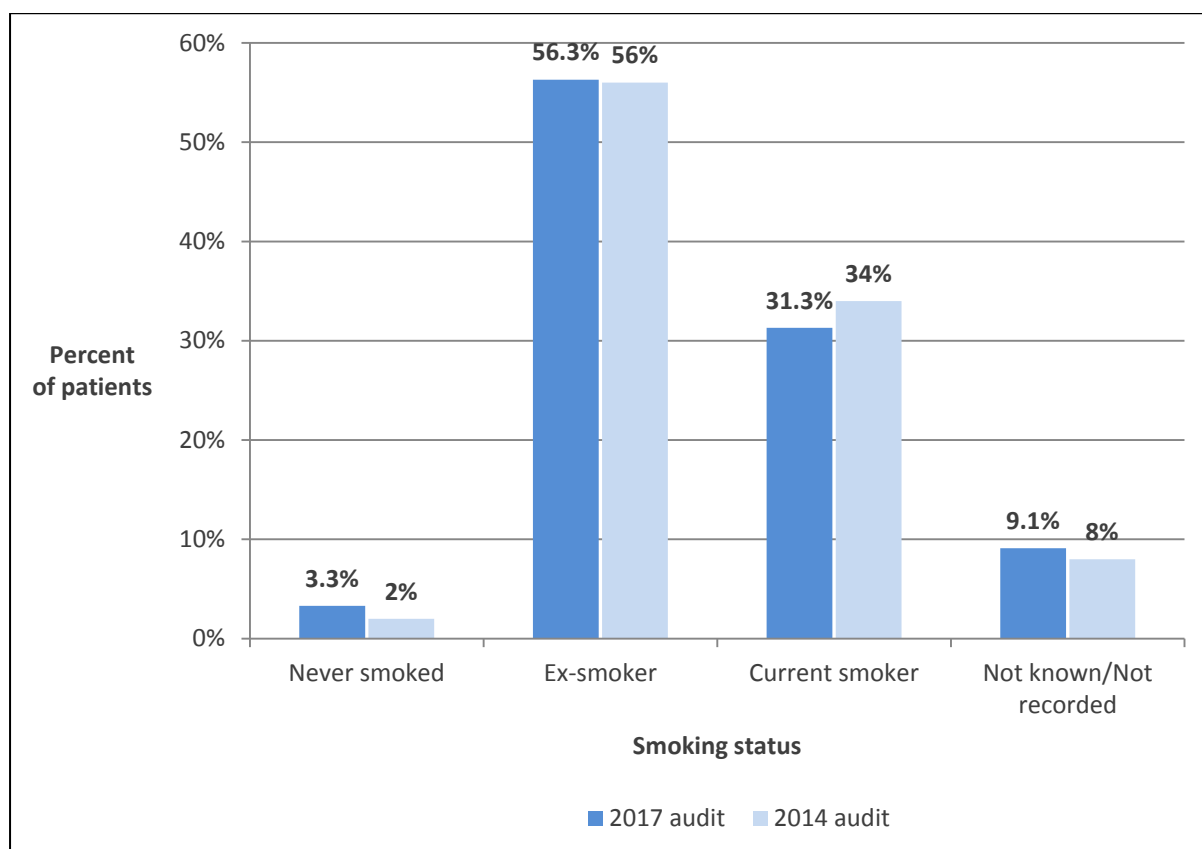
¹ Participants were asked in the dataset to record the value of the spirometric test: the patient's most recent FEV₁ as well as their most recent FVC. These have been used to calculate the FEV₁/FVC ratio (ie degree of airflow obstruction)

3.3. Smoking cessation

3.3.1 What was the smoking status for this patient, as documented for the current admission?

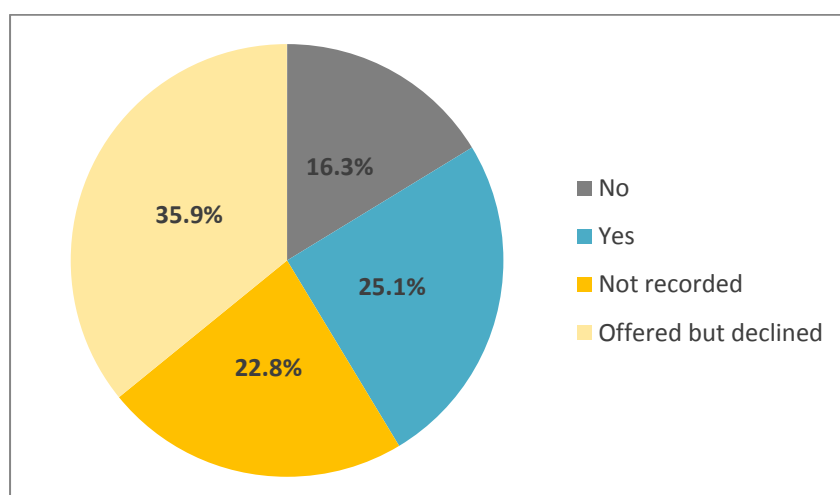
| Smoking status | 2017 (N=36,341) | 2014 (N=13,414) |
|------------------------|-----------------|-----------------|
| Never smoked | 3.3% (1,213) | 2% (310) |
| Ex-smoker | 56.3% (20,466) | 56% (7,552) |
| Current smoker | 31.3% (11,370) | 34% (4,528) |
| Not known/Not recorded | 9.1% (3,292) | 8% (1,024) |

Smoking status for patients included in the 2017 and 2014 audits



3.3.2 If a current smoker, was the patient prescribed smoking-cessation pharmacotherapy during the current admission?

| Prescribed smoking-cessation pharmacotherapy during the admission | 2017 (N=11,370) |
|---|-----------------|
| No | 16.3% (1,848) |
| Yes | 25.1% (2,855) |
| Not recorded | 22.8% (2,590) |
| Offered but declined | 35.9% (4,077) |

Current smokers that were prescribed smoking cessation pharmacotherapy during their admission**3.4. Dyspnoea, eosinopenia, consolidation, acidaemia and atrial fibrillation (DECAF) score****3.4.1 Was a DECAF score recorded for this patient?**

| DECAF score | 2017 (N=36,341) |
|-------------|-----------------|
| No | 83.5% (30,327) |
| Yes | 14.5% (5,278) |
| Not clear | 2.0% (736) |

3.4.2 If yes, what was the recorded DECAF score?

| DECAF score | 0 | 1 | 2 | 3 | 4 | 5 | 6 |
|---|------------------|------------------|------------------|----------------|---------------|--------------|----------------|
| 2017 (N=5,278) | 21.0% (1,109) | 33.0% (1,741) | 26.8% (1,412) | 14.1% (742) | 4.2% (224) | 0.9% (47) | 0.1% (3) |
| 2014^m (N=5,583) | 26% (1,449) | 41% (2,263) | 24% (1,342) | 8% (446) | 1% (79) | 0.1% (4) | Not applicable |

^m In the 2014 analysis a modified DECAF score was used as we could not distinguish between MRC dyspnoea grade 5a (score 1) and grade 5b (score 2) therefore a score of 1 was given to all MRC grade 5. For this reason the 2014 and 2017 scores should not be directly compared



Section 4: Non-invasive ventilation (NIV)

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Key findings

- The median time from arrival to acute treatment with NIV was 4.3 hours.

For the full key findings and recommendations, please see the national report COPD: Working together, available at www.rcplondon.ac.uk/working-together

Navigation

This section contains the following tables and graphs. If you are viewing this report on a computer, you can select the table that you wish to see from the list below.

- 4.1 NIV**
 - 4.1.1 Did the patient receive acute treatment with NIV?**
 - 4.1.2 If the patient received acute treatment with NIV, was it received within 3 hours of arrival?**
 - 4.1.3 If the patient received acute treatment with NIV was it received within 3 hours from admission? (2014 clinical audit figures)**
 - 4.1.4 Average time from arrival at hospital to acute treatment with NIV**
 - 4.1.5 Time from arrival to acute treatment with NIV**
 - 4.1.6 Time from admission to acute treatment with NIV (2014 clinical audit figures)**

4.1 NIV

4.1.1 Did the patient receive acute treatment with NIV?

| Acute treatment with NIV | 2017 (N=36,341) | 2014 (N=13,414) |
|----------------------------|-----------------|---------------------------|
| No | 89.1% (32,386) | 87% (11,600) ⁿ |
| Yes | 10.9% (3,955) | 12% (1,612) |
| Patient intubated directly | Not asked | 0.2% (30) |
| Not known | Not asked | 1% (172) |

4.1.2 If the patient received acute treatment with NIV, was it received within 3 hours of arrival?

| Acute treatment with NIV received within 3 hours of arrival | 2017 (N=3,955) |
|---|----------------|
| No | 44.7% (1,766) |
| Yes | 30.1% (1,191) |
| No time/date recorded | 25.2% (998) |

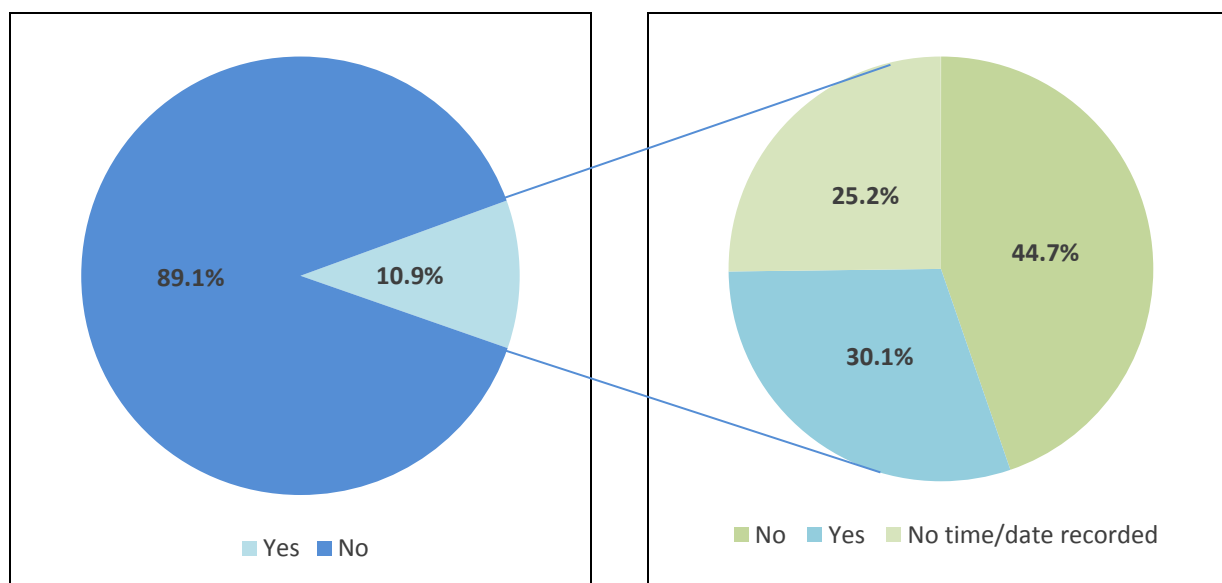
ⁿ Either not clinically indicated, patient declined, NIV not available or reason unclear

4.1.3 If the patient received acute treatment with NIV was it received within 3 hours from admission? (2014 clinical audit figures)^o

| Acute treatment with NIV received within 3 hours from admission | 2014 (N=1,191*) |
|---|-----------------|
| No | 57.6% (686) |
| Yes | 42.4% (505) |

*421 cases (1612 minus 1191) were excluded as either the date seen or the time of day, or both were not given

Patients in the 2017 audit who received NIV (left) within 3 hours (right)



4.1.4 Average time from arrival at hospital to acute treatment with NIV

| Time, in hours, from arrival at hospital to acute treatment with NIV | 2017 (N=2,956) |
|--|----------------|
| Mean | 19.7 |
| Standard deviation | 57.4 |
| Median | 4.3 |
| Interquartile range | 1.7–13.6 |

4.1.5 Time from arrival to acute treatment with NIV

| Time, in hours, from arrival at hospital to acute treatment with NIV | 2017 (N=2,956) |
|--|----------------|
| <3 hours | 40.3% (1,191) |
| 3–24 hours | 42.7% (1,263) |
| >24 hours | 17.0% (502) |

^o Results within table 4.1.3 have not been combined into table 4.1.2 as the metrics have been calculated and reported differently. In 2014 (4.1.3) it was NIV received within 3 hours of a patient's admission, whereas in 2017 (4.1.2) it was NIV received within 3 hours of a patient's arrival at hospital.

4.1.6 Time from admission to acute treatment with NIV (2014 clinical audit figures)^p

| Time, in hours, from admission at hospital to acute treatment with NIV | 2014 (N=1,096) |
|--|----------------|
| <3 hours | 37.4% (410) |
| 3–24 hours | 42.4% (465) |
| >24 hours | 20.2% (221) |

^p Results within table 4.1.6 have not been combined into table 4.1.5 as the metrics have been calculated and reported differently. In 2014 (4.1.6) it was time from admission to acute treatment with NIV where as in 2017 (4.1.5) it was time from arrival to acute treatment with NIV.



Section 5: Discharge processes

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Key findings

- Only **53%** of admissions received a **discharge bundle**. (*NICE [QS10] statement 8*)

For the full key findings and recommendations, please see the national report *COPD: Working together*, available at www.rcplondon.ac.uk/working-together

Navigation

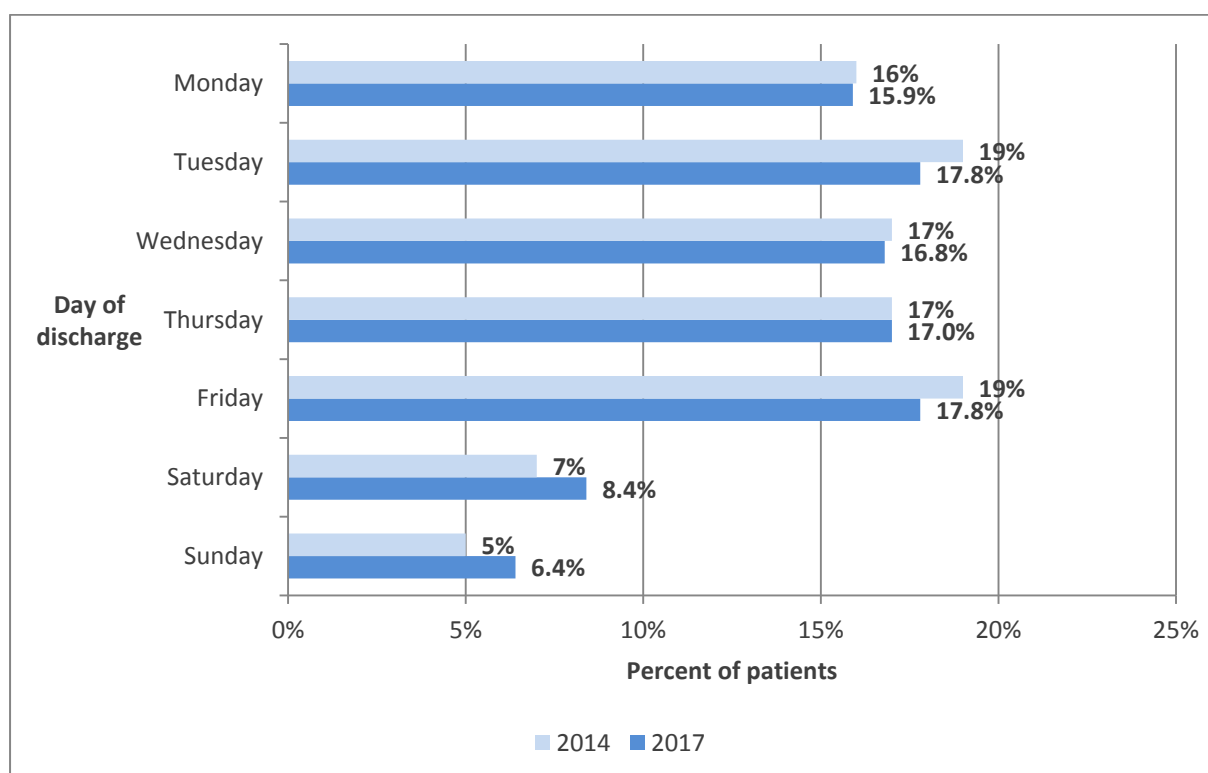
This section contains the following tables and graphs. If you are viewing this report on a computer, you can select the table that you wish to see from the list below.

- [5.1 Day of discharge](#)
 - [5.1.1 Average length of stay](#)
- [5.2 Did the patient die as an inpatient in your hospital?](#)
- [5.3 Has a British Thoracic Society \(BTS\), or equivalent, discharge bundle been completed for this admission?](#)
- [5.4 What follow-up arrangements have been made for this patient?](#)

5.1 Day of discharge

| | Day of discharge | | | | | | |
|----------------------------------|------------------|------------------|------------------|------------------|------------------|-----------------|-----------------|
| | Monday | Tuesday | Wednesday | Thursday | Friday | Saturday | Sunday |
| 2017 (N=36,341) | 15.9% (5,759) | 17.8% (6,472) | 16.8% (6,086) | 17.0% (6,175) | 17.8% (6,476) | 8.4% (3,043) | 6.4% (2,330) |
| 2014 (N=12,838) | 16% (2,090) | 19% (2,390) | 17% (2,180) | 17% (2,193) | 19% (2,384) | 7% (905) | 5% (674) |

Day of patient discharge in 2017 and 2014



5.1.1 Average length of stay

| Length of stay, days | 2017 (N=36,341) | 2014 (N=12,838) |
|----------------------|-----------------|-----------------|
| Mean | 5.8 | Not reported |
| Standard deviation | 7.8 | Not reported |
| Median | 4 | 4 |
| Interquartile range | 2–7 | 2–8 |

5.2 Did the patient die as an inpatient in your hospital?

| Inpatient mortality | 2017 (N=36,341) | 2014 (N=13,414) |
|---------------------|-----------------|-----------------|
| No | 96.1% (34,926) | 95.7% (12,838) |
| Yes | 3.9% (1,415) | 4.3% (576) |

5.3 Has a British Thoracic Society (BTS), or equivalent, discharge bundle been completed for this admission?

| Discharge bundle completed for this admission | 2017 (N=36,341) |
|---|-----------------|
| No | 35.8% (13,008) |
| Yes | 53.0% (19,275) |
| Not clear | 4.7% (1,719) |
| Patient self-discharged | 0.6% (213) |
| Patient died | 3.9% (1,415) |
| Other | 2.0% (711) |

5.4 What follow-up arrangements have been made for this patient?

| | 2017 (N=34,926) |
|--|-----------------|
| No arrangements apparent | 18.8% (6,575) |
| Patient discharged under the care of an early/assisted discharge team or integrated care service | 15.1% (5,276) |
| GP follow-up advised | 16.6% (5,800) |
| GP follow-up arranged | 2.7% (942) |
| Follow-up phone call scheduled | 8.0% (2,786) |
| Community respiratory clinic follow-up advised | 5.8% (2,016) |
| Community respiratory clinic follow-up arranged | 14.4% (5,042) |
| Hospital respiratory clinic follow-up advised | 5.9% (2,075) |
| Hospital respiratory clinic follow-up arranged | 24.7% (8,614) |
| No follow-up arranged, as discharged to other healthcare facility | 2.3% (787) |
| Discharged for end-of-life care | 0.7% (249) |
| Other | 12.5% (4,374) |



Section 6: Sub-analyses^q

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This section contains sub-analyses of the 2017 data presented in the previous sections, and presents associations between various metrics and:

- time from arrival to acute treatment with NIV
- review by a member of the respiratory team
- weekend admission.

Navigation

This section contains the following tables and graphs. If you are viewing this report on a computer, you can select the table that you wish to see from the list below.

- [6.1 Associations with time from arrival to acute treatment with NIV](#)
- [6.2 Associations with review by a member of the respiratory team within 24 hours](#)
- [6.3 Associations with weekend admission](#)

6.1 Associations with time from arrival to acute treatment with NIV

| | Time from arrival to acute treatment with NIV | | | p-value ^f |
|------------------------|---|-------------------------|----------------------|----------------------|
| | <3 hours (N=1,191) | 3–24 hours (N=1,263) | >24 hours (N=502) | |
| Length of stay >4 days | 67.3% (802) | 69.4% (871) | 84.5% (424) | <0.0001 |
| Died as inpatient | 10.4% (124) | 10.9% (137) | 24.7% (124) | <0.0001 |

Relative to patients who received acute treatment with NIV in under 3 hours:

- Patients who received **acute treatment with NIV more than 24 hours after arrival** were **2.6 times more likely** (odds ratio (OR): 2.63 [95% confidence interval (CI): 2.00–3.47]) to have a **length of stay greater** than the **median of 4 days** (p<0.0001). **It should be noted that reverse causation is likely an issue with this query as patients that need to stay longer may end up requiring NIV later.**
- Patients who received **acute treatment with NIV more than 24 hours after arrival** were **2.8 times more likely** (OR: 2.82 [95% CI: 2.13–3.74]) **to die as an inpatient** (p<0.0001).
- Patients who received acute treatment with NIV between 3 and 24 hours from arrival were 10% more likely (OR: 1.10 [95% CI: 0.93–1.30]) to have a length of stay greater than the median of 4 days (p=0.2822).
- Patients who received acute treatment with NIV between 3 and 24 hours from arrival were 6% more likely (OR: 1.06 [95% CI: 0.82–1.36]) to die as an inpatient (p=0.6798).

6.2 Associations with review by a member of the respiratory team within 24 hours

| | Specialist review within 24 hours | | p-value ^s |
|--------------------------------------|-----------------------------------|---------------|----------------------|
| | No | Yes | |
| Length of stay >4 days (N=36,341) | 41.7% (6,846) | 41.9% (8,355) | 0.673 |
| Died as an inpatient (N=36,341) | 4.3% (706) | 3.6% (709) | <0.001 |

^q Due to all outcomes other than mortality being quite common, odds ratios will not approximate risk ratios

^f χ^2 test for trend

^s χ^2 test

| | | | |
|--|---------------|----------------|--------|
| Oxygen prescribed (N=29,910) | 64.0% (8,496) | 74.1% (12,333) | <0.001 |
| Received NIV within 3 hours (N=3,955) | 25.9% (324) | 32.1% (867) | <0.001 |
| Received smoking cessation pharmacotherapy (N=8,780) | 69.4% (2,257) | 84.6% (4,675) | <0.001 |
| Received a discharge bundle (N=34,002) | 37.6% (5,748) | 72.3% (13,527) | <0.001 |

- Patients who **received a respiratory team review within 24 hours** (relative to those who didn't) were **18% less likely** (OR: 0.82 [95% CI: 0.74–0.91]) **to die as an inpatient**.
- Patients who received a respiratory team review within 24 hours and required oxygen were **61% more likely** (OR: 1.61 [95% CI: 1.53–1.70]) **to receive an oxygen prescription** than patients who did not receive a respiratory team review within 24 hours and required oxygen.
- Patients who received a respiratory team review within 24 hours and required NIV during their admission were **35% more likely** (OR: 1.35 [95% CI: 1.16–1.57]) **to receive NIV within 3 hours** than patients who did not receive a respiratory team review within 24 hours and required NIV during their admission.
- Patients who received a respiratory team review within 24 hours and were a current smoker were **2.4 times more likely** (OR: 2.42 [95% CI: 2.17–2.69]) **to receive smoking cessation pharmacotherapy** than patients who did not receive a respiratory team review within 24 hours and were a current smoker.
- Patients who received a respiratory team review within 24 hours and were discharged were **4.3 times more likely** (OR: 4.32 [95% CI: 4.11–4.53]) **to receive a discharge bundle** than patients who did not receive a respiratory team review within 24 hours and were discharged.

6.3 Associations with weekend admission

| | Weekend admission | | p-value ^t |
|--|-------------------|---------------|----------------------|
| | No | Yes | |
| Length of stay > 4 days (N=36,341) | 42.9% (11,711) | 38.6% (3,490) | <0.001 |
| Died as an inpatient (N=36,341) | 3.8% (1,039) | 4.2% (376) | 0.139 |
| Oxygen prescribed (N=29,910) | 69.7% (15,622) | 69.5% (5,207) | 0.779 |
| Received NIV within 3 hours (N=3,955) | 29.6% (866) | 31.7% (325) | 0.214 |
| Received smoking cessation pharmacotherapy (N=8,780) | 79.1% (5,217) | 78.4% (1,715) | 0.450 |
| Received a discharge bundle (N=34,002) | 57% (14,669) | 55% (4,606) | <0.001 |

- Patients who were admitted at the weekend (relative to those admitted during the week) were **17% less likely** (OR: 0.83 [95% CI: 0.80–0.88]) to have a length of stay greater than the median of 4.
- Patients who were admitted at the weekend and were discharged were **11% less likely** (OR: 0.89 [95% CI: 0.85–0.94]) to receive a discharge bundle than patients who were admitted during the week and were discharged.

^t X² test



Section 7: Web-tool run charts

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During 2017 the National COPD Audit Programme undertook a phased release of run charts via the secondary care audit web-tool (www.nacap.org.uk). The charts were derived from the data entered by hospitals registered for the audit, and introduced to be used by clinical teams to help drive quality improvement by providing near real-time feedback on service quality and compliance with national care standards. The charts display hospital level data, benchmarked to the national average and are only available to be viewed by registered users of the web-tool.

The charts depicted that between February 2017 and February 2018 there was improvement in performance (notably so for the BPT items) across most of the key indicators:

| Key indicator | February 2017 | February 2018 | Increase or decrease |
|--|---------------|---------------|----------------------|
| Best practice tariff (BPT)³ (released March 2017) | | | |
| Patients receiving a review by a member of the respiratory team during admission | 72.7% | 91.2% | ↑ |
| Patients receiving a review by a member of the respiratory team within 24 hours of admission | 48.4% | 69.3% | ↑ |
| Patients receiving a discharge bundle upon discharge | 48.2% | 81.5% | ↑ |
| Patients where care meets the BPT for COPD | 31.1% | 62.0% | ↑ |
| Oxygen (released May 2017) | | | |
| Patients being prescribed oxygen | 70.3% | 74.7% | ↑ |
| Patients being prescribed oxygen to target saturation | 95.7% | 97.7% | ↑ |
| Spirometry (released May 2017) | | | |
| Patients with spirometry result available | 36.4% | 40.8% | ↑ |
| Smoking cessation pharmacotherapy (released June 2017) | | | |
| Current smokers that have been prescribed smoking cessation pharmacotherapy | 23.9% | 26.6% | ↑ |
| NIV (released July 2017) | | | |
| Patients receiving NIV | 11.8% | 10.1% | ↓ |
| Patients receiving NIV within 3 hours of arrival | 26.6% | 29.8% | ↑ |

Appendix A: Report preparation

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This report was written by the following people, on behalf of the national COPD secondary care audit 2017 workstream group (the full list of workstream group members is included in [Appendix G](#)).

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Appendix B: Introduction to the National COPD Audit Programme

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The National COPD Audit Programme is a programme of work that aims to drive improvements in the quality of care and services provided for patients with COPD in England and Wales. The programme looks at COPD care across the patient pathway, both in and out of hospital, bringing together key elements from the primary, secondary and community care sectors.

There are three programme workstreams.

1. Primary care: collection of audit data from general practice patient record systems in Wales. Delivered by the RCP and NHS Digital, working with the Primary Care Respiratory Society UK, the Royal College of General Practitioners and the NHS Wales Informatics Service. Another round of audit took place in 2017.
2. Secondary care: in 2014, there were snapshot audits of patients who were admitted to hospital with COPD exacerbation, plus organisational audits of the resourcing of COPD services in acute units. The 2014 audits were delivered by the BTS, working with the RCP. A continuous audit of admission to hospital with COPD exacerbation commenced in 2017.
3. Pulmonary rehabilitation: audits of COPD patients attending pulmonary rehabilitation (including outcomes at 180 days), plus organisational audits of the resourcing of pulmonary rehabilitation services for COPD patients. The 2015 round of this audit was delivered by the BTS, working with the RCP. Another round of snapshot clinical and organisational audits took place in 2017.

The audit also delivered a 1-year development project exploring the potential/feasibility of future incorporation of a patient-reported experience measure (PREM) into the audit programme. This was delivered by the British Lung Foundation, working with the Picker Institute Europe.

The programme is commissioned by HQIP as part of the National Clinical Audit (NCA) Programme. It is included in the list of national audits for inclusion in NHS trusts' quality accounts and also the NHS Wales Clinical Audit and Outcome Review Plan.

Appendix C: Methodology

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Methodology of the audit creation and setup

The National COPD Audit Programme's secondary care clinical audit 2017 built upon the learning from the 2014 clinical audit.^u The structure of the dataset was similar to that used in 2014, however, it was considerably streamlined to account for the change in methodology from snapshot (in 2014) to continuous (in 2017) audit.

All secondary care hospitals in England and Wales that admit patients with acute exacerbations of COPD (AECOPD) were approached to participate in the audit. Out of 197 eligible, 182 hospitals participated in the clinical audit. A full list of participating hospitals, including those hospitals that opted out of taking part are listed in [Appendix F](#).

The continuous audit launched on 1 February 2017. This report presents the results of the cohort of patients discharged between the audit's launch date and 13 September 2017.

Information governance

The audit involved the collection of patient identifiable data for the purpose of linkage with data from other sources (such as Hospital Episode Statistics and Office for National Statistics data for readmission and mortality data), and the audit operated with Section 251 approval via the Confidential Advisory Group (CAG), reference number: CAG 8-06(b)/2013.^v The rationale for this was the acuity of the patient cohort and the rate of admissions (approximately 115,000 per annum),^w meaning that the seeking of patient consent (the alternative route for collection of identifiable patient data) was not a viable option.

A patient leaflet and poster, fair processing information as well as the audit data flows, were made available on the audit website www.rcplondon.ac.uk/projects/outputs/secondary-care-audit-2017-resources.

Recruitment

There was a single recruitment process for both the clinical and organisational secondary care audits, which began in early 2016, using the following channels:

- partner and stakeholder channels (such as the BTS's e-bulletin, and the Association of Respiratory Nurse Specialists' newsletter)
- Twitter and the audit's own newsletter
- communication with hospitals that participated in the 2014 audit
- direct letters to chief executives and medical directors.

The reasons provided to participate in the audits were as follows:

^u Stone RA, Holzhauser-Barrie J, Lowe D *et al.* COPD: Who cares matters. National Chronic Obstructive Pulmonary Disease (COPD) Audit Programme: Clinical audit of COPD exacerbations admitted to acute units in England and Wales 2014. National clinical audit report. London: RCP, February 2015. www.rcplondon.ac.uk/projects/outputs/copd-who-cares-matters-clinical-audit-2014 [Accessed 16 February 2018].

^v A record of the audit approval can be found at www.hra.nhs.uk/about-the-hra/our-committees/section-251/cag-advice-and-approval-decisions/ (April 2013 onwards; non research) [Accessed 16 February 2018].

^w <https://statistics.blf.org.uk/copd> [Accessed 16 February 2018].

- the status of the audit as part of NHS Quality Accounts
- the use of the clinical audit to support the launch of the new BPT for COPD (launched on 1 April 2017)
- the Care Quality Commission's (CQC's) future use (as of autumn 2017) of clinical audit metrics in its hospital inspections
- the alignment of the audit to National Institute for Health and Care Excellence (NICE) guidance and key findings from the 2014 audit
- the fact the audit/s would prove to be a useful tool for facilitating local improvement.

Hospitals were asked to complete a registration form, nominating a 'lead clinician' and listing any other team members that would form part of the audit team. It was made clear to prospective participants that the 'lead clinician' role took ultimate responsibility for the accuracy of the data that were entered for that hospital.

Once hospitals had registered, they were sent a Caldicott Guardian letter and a form to have signed off. Only after the Caldicott Guardian form was received by the audit team at the RCP was the hospital considered 'fully signed up' and, at that point, it was registered on the audit web-tool.

People were registered on the web-tool as having one of two roles: 'lead clinician' or 'data inputters'. The former were able to approve the creation of new users for that hospital (ie as the audits progressed) and ensure that they were suitable from an information governance perspective.

The audit team chased the registration form and Caldicott Guardian form up until the audit start date.

Audit question development and pilot

The audit dataset was based on the equivalent 2014 dataset. It was developed iteratively by the audit programme team and clinical lead, in consultation with the workstream group, in particular the representatives of the BTS and the Society of Acute Medicine.

The dataset and web-tool were tested in 33 hospitals (highlighted in the list of participating hospitals in [Appendix F](#)) in a pilot between Monday 13 August and Friday 7 September 2016. The pilot hospitals were asked to feed back on the web-tool, the audit questions and the help notes provided.

This feedback was discussed by the team and workstream group, and the dataset was finalised. It is available to download from our website: www.rcplondon.ac.uk/projects/outputs/secondary-care-audit-2017-resources.

The audit team created a full dataset containing helpnotes and reasons for the inclusion of each data item, and also a two-sided proforma to aid those hospitals who prefer to collect the information on paper.

Soft launch period

To aid local planning, the audit team made the clinical audit available on the web-tool from Tuesday 1 November 2016) in 'soft launch' form. Users were able to:

- access the full dataset
- review the help notes
- input data and test the validation rules.

They were, however, **not able to save results**. This gave the teams approximately 3 months to prepare, prior to the audit's launch at the start of February. This was felt to be particularly important, as the dataset would need to be completed multiple times by participants, and familiarisation with the data items required would be key.

Data entry

Hospitals were required to enter data via the audit programme's bespoke web-tool, created by Crown Informatics Ltd (available at www.nacap.org.uk).

Documentation to support participation in the audit, including audit instructions, data collection sheets, datasets with help notes, as well as copies of all communications and newsletters, was available on the secondary care audit resources website and on the web-tool.

Regular email updates and newsletters were sent to participants throughout the data collection period, with reminders about timelines and answers to frequently asked questions.

Data storage, security and transfer

Data were collected on the audit's bespoke web-tool. These data were stored and processed at a secure data centre, owned by AIMES Grid Services, located in Liverpool, UK. It operates to ISO 27001 certification (2015). The servers are owned and operated by Crown Informatics Ltd and are held in a secure locked rack, accessible to named individuals. All access is logged, managed and supervised.

This data centre provides N3 aggregation in collaboration with NHS Digital. Data are stored in secured databases (software by IBM) and encrypted on disc (AES256 standard) and additionally in the database where required. Backups are encrypted at AES256, held in dual copies and stored securely.

Crown Informatics Ltd operate secure SSL at 256 bit, using SHA256 (SHA2) signatures and 4096-bit certificates. Crown Informatics Ltd's certificate is an 'OV' certified by a respected global certifier (Starfield/GoDaddy). In addition, 'Qualsys' using 'SSL Labs' have given the audit site an 'A' rating.

At the end of the data collection period, the data were extracted from the web-tool by the central audit team, using an 'extract' provision developed by Crown Informatics. They were then transferred securely (using the RCP's Mimecast system) to the team at Imperial College London for analysis.

Telephone and email support

The audit programme team at the RCP provided a helpdesk every working day during office hours, which was available via both telephone and email, so that participants could contact the team directly with any questions.

Analysis methodology

The data were analysed at Imperial College London (National Heart and Lung Institute). Data received from the RCP were imported in to Stata 15.

A few patients with unlikely ages (less than 35 years of age) and negative lengths of stay were discovered so these records were sent back to hospitals via the RCP for querying. Once correct results were received back, the appropriate corrections were made to the dataset.

The dataset was then prepared, cleaned, and analysed as follows:

- All string categorical variables were recoded numerically and labelled with the former string value.
- All string date/time variables were converted to numerical date/time variables.
- All indicator variables (to denote presence or absence) were converted from their current format (eg an 'X' character) to a binary 0 or 1 value.
- Variables that comprised lists of strings were converted to separate binary variables for each possible string in the list to indicate the presence or absence of each possible item.
- All draft entries were removed (n=860).
- Admissions before arrival were removed (n=0).
- Admissions after discharge were removed (n=2).
- Respiratory specialist reviews before arrival were removed (n=0).
- Respiratory specialist reviews after discharge were removed (n=224).
- NIVs before arrival were removed (n=95).
- NIVs after discharge were removed (n=34).
- Discharges before arrival were removed (n=0).
- English and Welsh quintiles of index of multiple deprivation (IMD) were produced (1=most deprived, 5=least deprived) using the provided IMD rank for each patient.
- Number of admissions for each hospital was generated.
- Two-hour time categories (ie 12 total) were generated for time of arrival and time of first review by a member of the respiratory team.
- Time from arrival to admission was generated by subtracting arrival time from admission time and patients with admission wait times greater than or equal to 24 hours were removed as this was considered unrealistic (n=387).
- Time from admission to specialist review was generated by subtracting admission time from review time and patients with review wait times less than or equal to -24 hours (24 hours prior) were removed as this was considered unrealistic (n=0 [removed in previous stage]).
- Time from arrival to NIV was calculated by subtracting arrival time from time of NIV administration.
- FEV₁/FVC ratio was cleaned such that values under 0.2 or over 1.0 were replaced with missing values and a variable to denote airflow obstruction for anyone with a ratio under 0.7 was created.
- Errors in day of week variables were corrected using Stata 'dow()' command to get the correct day of the week for the specified date (n=1).
- Errors in the length of stay variable were corrected by subtracting date of admission from date of discharge (n=4).
- Variables required for analysis and generating odds ratios were created:
 - Specialist review in 24 hours (yes (y) / no (n))
 - Oxygen prescription (y/n)
 - NIV in 3 hours (y/n)
 - Time to NIV administration (<3 hours, 3–24 hours, >24 hours)
 - Smoking cessation pharmacotherapy prescribed (y/n)
 - Discharge bundle completed (y/n)
 - Weekend admission (y/n)
 - Length of stay (equal to or below median/above median).
- All Stata analysis was performed using the 'summarize', 'tabulate', or 'tabodds' commands, depending on whether median/mean, proportion, or odds ratio was the desired analysis, respectively.

Appendix D: Chronic obstructive pulmonary disease in adults NICE Quality standard [QS10]

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Please note: In 2016 this quality standard was updated and statements prioritised in 2011 were updated (2011, updated 2016) or replaced (new 2016).

Statements are marked as [new 2016] or [2011, updated 2016]:

- [new 2016] if the statement covers a new area for quality improvement
- [2011, updated 2016] if the statement covers an area for quality improvement included in the 2011 quality standard and has been updated.

| No. | Quality statement |
|-----|---|
| 1 | People aged over 35 years who present with a risk factor and one or more symptoms of chronic obstructive pulmonary disease (COPD) have post-bronchodilator spirometry. [2011, updated 2016] |
| 2 | People with COPD who are prescribed an inhaler have their inhaler technique assessed when starting treatment and then regularly during treatment. [2011, updated 2016] |
| 3 | People with stable COPD and a persistent resting stable oxygen saturation level of 92% or less have their arterial blood gases measured to assess whether they need long-term oxygen therapy. [2011, updated 2016] |
| 4 | People with stable COPD and exercise limitation due to breathlessness are referred to a pulmonary rehabilitation programme. [2011, updated 2016] |
| 5 | People admitted to hospital for an acute exacerbation of COPD start a pulmonary rehabilitation programme within 4 weeks of discharge. [2011, updated 2016] |
| 6 | People receiving emergency oxygen for an acute exacerbation of COPD have oxygen saturation levels maintained between 88% and 92%. [new 2016] |
| 7 | People with an acute exacerbation of COPD and persistent acidotic hypercapnic ventilatory failure that is not improving after 1 hour of optimal medical therapy have non-invasive ventilation. [2011, updated 2016] |
| 8 | (Placeholder ^x) Hospital discharge care bundle. [new 2016] |

^x A placeholder statement is an area of care that has been prioritised by the Quality Standards Advisory Committee but for which no source guidance is currently available. A placeholder statement indicates the need for evidence-based guidance to be developed in this area.

Statements from the 2011 quality standard for COPD that may still be useful at a local level, but are no longer considered national priorities for improvement:

- People with COPD have a current individualised comprehensive management plan, which includes high-quality information and educational material about the condition and its management, relevant to the stage of disease.
- People with COPD have a comprehensive clinical and psychosocial assessment, at least once a year or more frequently if indicated, which includes degree of breathlessness, frequency of exacerbations, validated measures of health status and prognosis, presence of hypoxaemia and comorbidities.
- People with COPD who smoke are regularly encouraged to stop and are offered the full range of evidence-based smoking cessation support.
- People who have had an exacerbation of COPD are provided with individualised written advice on early recognition of future exacerbations, management strategies (including appropriate provision of antibiotics and corticosteroids for self-treatment at home) and a named contact.
- People with COPD receiving long-term oxygen therapy are reviewed in accordance with NICE guidance, at least annually, by a specialist oxygen service as part of the integrated clinical management of their COPD.
- People admitted to hospital with an exacerbation of COPD are cared for by a respiratory team, and have access to a specialist early supported-discharge scheme with appropriate community support.
- People admitted to hospital with an exacerbation of COPD are reviewed within 2 weeks of discharge.
- People with advanced COPD, and their carers, are identified and offered palliative care that addresses physical, social and emotional needs.

Appendix E: Chronic obstructive pulmonary disease in over 16s: diagnosis and management NICE Clinical guideline [CG101]

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Below is only a summary of the NICE Clinical guidelines [CG101] that we specifically refer to within this report. To see the full guideline please use the following link: www.nice.org.uk/guidance/cg101

| | |
|--------------|---|
| 1.1 | Diagnosing COPD |
| 1.1.8 | Referral for specialist advice |
| 1.1.8.2 | Patients who are referred do not always have to be seen by a respiratory physician. In some cases they may be seen by members of the COPD team who have appropriate training and expertise. [2004] |
| 1.1.2 | Spirometry |
| 1.1.2.1 | Spirometry should be performed: <ul style="list-style-type: none"> • at the time of diagnosis • to reconsider the diagnosis, if patients show an exceptionally good response to treatment. [2004] |
| 1.1.2.2 | Measure post-bronchodilator spirometry to confirm the diagnosis of COPD. [new 2010] |
| 1.1.2.3 | Consider alternative diagnoses or investigations in: <ul style="list-style-type: none"> • older people without typical symptoms of COPD where the FEV₁/FVC ratio is <0.7 • younger people with symptoms of COPD where the FEV₁/FVC ratio is ≥0.7. [new 2010] |
| 1.1.2.4 | All health professionals involved in the care of people with COPD should have access to spirometry and be competent in the interpretation of the results. [2004] |
| 1.1.2.5 | Spirometry can be performed by any healthcare worker who has undergone appropriate training and who keeps his or her skills up to date. [2004] |
| 1.1.2.6 | Spirometry services should be supported by quality control processes. [2004] |
| 1.1.2.7 | It is recommended that ERS 1993 reference values ^y are used but it is recognised that these values may lead to under-diagnosis in older people and are not applicable in black and Asian populations. ^z [2004] |

| | |
|--------------|--|
| 1.2 | Managing stable COPD |
| 1.2.1 | Smoking cessation |
| 1.2.1.1 | An up-to-date smoking history, including pack years smoked (number of cigarettes smoked per day, divided by 20, multiplied by the number of years smoked), should be documented for everyone with COPD. [2004] |
| 1.2.1.2 | All COPD patients still smoking, regardless of age, should be encouraged to stop, and offered help to do so, at every opportunity. [2004] |
| 1.2.1.3 | Unless contraindicated, offer NRT, varenicline or bupropion, as appropriate, to people who are planning to stop smoking combined with an appropriate support programme to optimise smoking quit rates for people with COPD. ^{aa} [2010] |
| 1.2.6 | Non-invasive ventilation |
| 1.2.6.1 | Adequately treated patients with chronic hypercapnic respiratory failure who have |

^y Quanjer PH, Tammeling GJ, Cotes JE *et al*. Lung volumes and forced ventilatory flows. Report Working Party Standardization of Lung Function Tests, European Community for Steel and Coal. Official Statement of the European Respiratory Society. *Eur Respir J Suppl* 1993;16;5–40.

^z Definitive spirometry reference values are not currently available for all ethnic populations. The GDG was aware of ongoing research in this area.

^{aa} Stop smoking services. NICE Public health guidance [PH10]. www.nice.org.uk/guidance/ph10 [Accessed January 2018].

| | |
|--|---|
| | required assisted ventilation (whether invasive or non-invasive) during an exacerbation or who are hypercapnic or acidotic on LTOT should be referred to a specialist centre for consideration of long-term NIV. [2004] |
|--|---|

| | |
|---------------|---|
| 1.3 | Management of exacerbations of COPD |
| 1.3.6 | Oxygen therapy during exacerbations of COPD^{bb} |
| 1.3.6.1 | The oxygen saturation should be measured in patients with an exacerbation of COPD, if there are no facilities to measure arterial blood gases. [2004] |
| 1.3.6.2 | If necessary, oxygen should be given to keep the SaO ₂ within the individualised target range. [2004, amended 2010] |
| 1.3.6.3 | Pulse oximeters should be available to all healthcare professionals involved in the care of patients with exacerbations of COPD and they should be trained in their use. Clinicians should be aware that pulse oximetry gives no information about the PCO ₂ or pH. [2004] |
| 1.3.6.4 | When the patient arrives at hospital, arterial blood gases should be measured and the inspired oxygen concentration noted in all patients with an exacerbation of COPD. Arterial blood gas measurements should be repeated regularly, according to the response to treatment. [2004] |
| 1.3.7 | Non-invasive ventilation (NIV) and COPD exacerbations |
| 1.3.7.1 | NIV should be used as the treatment of choice for persistent hypercapnic ventilatory failure during exacerbations despite optimal medical therapy. [2004] |
| 1.3.7.2 | It is recommended that NIV should be delivered in a dedicated setting with staff who have been trained in its application, who are experienced in its use and who are aware of its limitations. [2004] |
| 1.3.7.3 | When patients are started on NIV there should be a clear plan covering what to do in the event of deterioration and ceilings of therapy should be agreed. [2004] |
| 1.3.11 | Discharge planning |
| 1.3.11.1 | Spirometry should be measured in all patients before discharge. [2004] |
| 1.3.11.2 | Patients should be re-established on their optimal maintenance bronchodilator therapy before discharge. [2004] |
| 1.3.11.3 | Patients who have had an episode of respiratory failure should have satisfactory oximetry or arterial blood gas results before discharge. [2004] |
| 1.3.11.4 | All aspects of the routine care that patients receive (including appropriateness and risk of side effects) should be assessed before discharge. [2004] |
| 1.3.11.5 | Patients (or home carers) should be given appropriate information to enable them to fully understand the correct use of medications, including oxygen, before discharge. [2004] |
| 1.3.11.6 | Arrangements for follow-up and home care (such as visiting nurse, oxygen delivery, referral for other support) should be made before discharge. [2004] |
| 1.3.11.7 | Before the patient is discharged, the patient, family and physician should be confident that he or she can manage successfully. When there is remaining doubt a formal activities of daily living assessment may be helpful. [2004] |

^{bb} The exacerbation section of this guideline was outside the scope of the 2010 update. However the GDG was aware that some recommendations in this section of the guideline were out of date, and these have been removed. Readers should refer to local protocols.

Appendix F: Participating hospitals in the 2017 clinical secondary care audit

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Participating NHS acute units in England:

| Trust | Unit |
|---|-------------------------------------|
| Aintree University Hospitals NHS Foundation Trust | Aintree University Hospital |
| Airedale NHS Foundation Trust | Airedale General Hospital |
| Ashford and St Peter's Hospital NHS Foundation Trust | St Peter's Hospital |
| Barking Havering and Redbridge University Hospitals NHS Trust | King George Hospital |
| Barnsley Hospital NHS Foundation Trust | Barnsley Hospital |
| Barts Health NHS Trust | Newham University Hospital |
| Barts Health NHS Trust | The Royal London Hospital |
| Basildon and Thurrock University Hospitals NHS Foundation Trust | Basildon Hospital |
| Bedford Hospital NHS Trust | Bedford Hospital |
| Blackpool Teaching Hospitals NHS Foundation Trust | Blackpool Victoria Hospital |
| Bolton NHS Foundation Trust | Royal Bolton Hospital |
| Bradford Teaching Hospitals NHS Foundation Trust | Bradford Royal Infirmary |
| Brighton and Sussex University Hospitals NHS Trust | Princess Royal Hospital |
| Brighton and Sussex University Hospitals NHS Trust | Royal Sussex County Hospital |
| Buckinghamshire Healthcare NHS Trust | Stoke Mandeville Hospital |
| Burton Hospitals NHS Foundation Trust | Queen's Hospital |
| Calderdale and Huddersfield NHS Foundation Trust | Calderdale Royal Hospital |
| Calderdale and Huddersfield NHS Foundation Trust | Huddersfield Royal Infirmary |
| Central Manchester University Hospitals NHS Foundation Trust | Manchester Royal Infirmary |
| Central Manchester University Hospitals NHS Foundation Trust | Trafford General Hospital |
| Chelsea and Westminster Hospital NHS Foundation Trust | Chelsea and Westminster Hospital |
| Chelsea and Westminster Hospital NHS Foundation Trust | West Middlesex University Hospital |
| Chesterfield Royal Hospital NHS Foundation Trust | Chesterfield Royal Hospital |
| City Hospitals Sunderland NHS Foundation Trust | Sunderland Royal Hospital |
| Colchester Hospital University NHS Foundation Trust | Colchester General Hospital |
| Countess of Chester Hospital NHS Foundation Trust | Countess of Chester Hospital |
| County Durham and Darlington NHS Foundation Trust | Darlington Memorial Hospital |
| County Durham and Darlington NHS Foundation Trust | University Hospital of North Durham |
| Croydon Health Services NHS Trust | Croydon University Hospital |
| Dartford and Gravesham NHS Trust | Darent Valley Hospital |
| Derby Teaching Hospitals NHS Foundation Trust | Royal Derby Hospital |
| Doncaster and Bassetlaw Teaching Hospitals NHS Foundation Trust | Bassetlaw Hospital |
| Doncaster and Bassetlaw Teaching Hospitals NHS Foundation Trust | Doncaster Royal Infirmary |
| Dorset County Hospital NHS Foundation Trust | Dorset County Hospital |

| Trust | Unit |
|---|---|
| East and North Hertfordshire NHS Trust | Lister Hospital |
| East Cheshire NHS Trust | Macclesfield District General Hospital |
| East Kent Hospitals University NHS Foundation Trust | Kent and Canterbury Hospital |
| East Kent Hospitals University NHS Foundation Trust | Queen Elizabeth the Queen Mother Hospital |
| East Kent Hospitals University NHS Foundation Trust | William Harvey Hospital |
| East Lancashire Hospitals NHS Trust | Royal Blackburn Hospital |
| East Sussex Healthcare NHS Trust | Conquest Hospital |
| East Sussex Healthcare NHS Trust | Eastbourne District General Hospital |
| Epsom and St Helier University Hospitals NHS Trust | Epsom Hospital |
| Epsom and St Helier University Hospitals NHS Trust | St Helier Hospital |
| Frimley Health NHS Foundation Trust | Frimley Park Hospital |
| Frimley Health NHS Foundation Trust | Wexham Park Hospital |
| George Eliot Hospital NHS Trust | George Eliot Hospital |
| Gloucestershire Hospitals NHS Foundation Trust | Cheltenham General Hospital |
| Gloucestershire Hospitals NHS Foundation Trust | Gloucestershire Royal Hospital |
| Great Western Hospitals NHS Foundation Trust | The Great Western Hospital |
| Guy's and St Thomas' NHS Foundation Trust | St Thomas' Hospitals |
| Hampshire Hospitals NHS Foundation Trust | Basingstoke and North Hampshire Hospital |
| Hampshire Hospitals NHS Foundation Trust | Royal Hampshire County Hospital |
| Harrogate and District NHS Foundation Trust | Harrogate District Hospital |
| Heart of England NHS Foundation Trust | Birmingham Heartlands Hospital |
| Heart of England NHS Foundation Trust | Good Hope Hospital |
| Heart of England NHS Foundation Trust | Solihull Hospital |
| Hinchingbrooke Health Care NHS Trust | Hinchingbrooke Hospital |
| Homerton University Hospital NHS Foundation Trust | Homerton University Hospital |
| Hull and East Yorkshire Hospitals NHS Trust | Hull Royal Infirmary |
| Imperial College Healthcare NHS Trust | Charing Cross Hospital |
| Imperial College Healthcare NHS Trust | St Mary's Hospital |
| Ipswich Hospital NHS Trust | Ipswich Hospital |
| Isle of Wight NHS Trust | St Mary's Hospital |
| James Paget University Hospitals NHS Foundation Trust | James Paget Hospital |
| Kettering General Hospital NHS Foundation Trust | Kettering General Hospital |
| King's College Hospital NHS Foundation Trust | King's College Hospital |
| King's Lynn NHS Foundation Trust | The Queen Elizabeth Hospital |
| Kingston Hospital NHS Foundation Trust | Kingston Hospital |
| Lancashire Teaching Hospitals NHS Foundation Trust | Chorley and South Ribble Hospital |
| Lancashire Teaching Hospitals NHS Foundation Trust | Royal Preston Hospital |
| Lewisham and Greenwich NHS Trust | University Hospital Lewisham |
| London North West Healthcare NHS Trust | Ealing Hospital |
| London North West Healthcare NHS Trust | Northwick Park Hospital |
| Luton and Dunstable Hospital NHS Foundation Trust | Luton and Dunstable Hospital |
| Maidstone and Tunbridge Wells NHS Trust | Maidstone Hospital |
| Maidstone and Tunbridge Wells NHS Trust | Tunbridge Wells Hospital |

| Trust | Unit |
|---|--|
| Medway NHS Foundation Trust | Medway Maritime Hospital |
| Mid Cheshire Hospitals NHS Foundation Trust | Leighton Hospital |
| Mid Essex Hospital Services NHS Trust | Broomfield Hospital |
| Milton Keynes University Hospital NHS Foundation Trust | Milton Keynes Hospital |
| Norfolk and Norwich University Hospitals NHS Foundation Trust | Norfolk and Norwich University Hospital |
| North Bristol NHS Trust | Southmead Hospital |
| North Middlesex University Hospital NHS Trust | North Middlesex University Hospital |
| North Tees and Hartlepool NHS Foundation Trust | University Hospital of North Tees |
| Northampton General Hospital NHS Trust | Northampton General Hospital |
| Northern Devon Healthcare NHS Trust | North Devon District Hospital |
| Northern Lincolnshire and Goole NHS Foundation Trust | Diana Princess of Wales Hospital |
| Northern Lincolnshire and Goole NHS Foundation Trust | Scunthorpe General Hospital |
| Northumbria Healthcare NHS Foundation Trust | Northumbria Specialist Emergency Care Hospital |
| Nottingham University Hospitals NHS Trust | Nottingham City Hospital and Queen's Medical Centre Nottingham |
| Oxford University Hospitals NHS Trust | John Radcliffe Hospital |
| Pennine Acute Hospitals NHS Trust | Fairfield General Hospital |
| Pennine Acute Hospitals NHS Trust | North Manchester General Hospital |
| Pennine Acute Hospitals NHS Trust | Rochdale Infirmary |
| Peterborough and Stamford Hospitals NHS Foundation Trust | Peterborough City Hospital |
| Poole Hospital NHS Foundation Trust | Poole Hospital |
| Portsmouth Hospitals NHS Trust | Queen Alexandra Hospital |
| Royal Berkshire NHS Foundation Trust | Royal Berkshire Hospital |
| Royal Cornwall Hospitals NHS Trust | Royal Cornwall Hospital |
| Royal Devon and Exeter NHS Foundation Trust | Royal Devon and Exeter NHS Hospital |
| Royal Free London NHS Foundation Trust | Barnet Hospital |
| Royal Free London NHS Foundation Trust | Royal Free Hospital |
| Royal Liverpool and Broadgreen University Hospitals NHS Trust | Royal Liverpool University Hospital |
| Royal Surrey County Hospital NHS Foundation Trust | Royal Surrey County Hospital |
| Royal United Hospitals Bath NHS Foundation Trust | Royal United Hospital Bath |
| Salford Royal NHS Foundation Trust | Salford Royal Hospital |
| Salisbury NHS Foundation Trust | Salisbury District Hospital |
| Sandwell and West Birmingham Hospitals NHS Trust | City Hospital |
| Sandwell and West Birmingham Hospitals NHS Trust | Sandwell General Hospital |
| Sheffield Teaching Hospital NHS Foundation Trust | Northern General Hospital |
| Sherwood Forest Hospitals NHS Foundation Trust | Kings Mill Hospital |
| South Tees Hospitals NHS Foundation Trust | Friarage Hospital |
| South Tees Hospitals NHS Foundation Trust | The James Cook University Hospital |
| South Tyneside NHS Foundation Trust | South Tyneside District Hospital |
| South Warwickshire NHS Foundation Trust | Warwick Hospital |
| Southend University Hospital NHS Foundation Trust | Southend University Hospital |
| Southern Health NHS Foundation Trust | Lymington New Forest Hospital |

| Trust | Unit |
|---|--|
| Southport and Ormskirk Hospitals NHS Trust | Southport and Formby District General Hospital |
| St George's University Hospitals NHS Foundation Trust | St George's Hospital |
| St Helens and Knowsley Teaching Hospital NHS Trust | Whiston Hospital |
| Stockport NHS Foundation Trust | Stepping Hill Hospital |
| Surrey and Sussex Healthcare NHS Trust | East Surrey Hospital |
| Tameside and Glossop Integrated Care NHS Foundation Trust | Tameside General Hospital |
| Taunton and Somerset NHS Foundation Trust | Musgrove Park Hospital |
| The Dudley Group of Hospitals NHS Foundation Trust | Russells Hall Hospital |
| The Hillingdon Hospitals NHS Foundation Trust | Hillingdon Hospital |
| The Mid Yorkshire Hospitals NHS Trust | Dewsbury District Hospital |
| The Mid Yorkshire Hospitals NHS Trust | Pinderfields General Hospital |
| The Newcastle upon Tyne Hospitals NHS Foundation Trust | Royal Victoria Infirmary |
| The Princess Alexandra NHS Hospital Trust | Princess Alexandra Hospital |
| The Rotherham NHS Foundation Trust | Rotherham General Hospital |
| The Royal Bournemouth and Christchurch Hospitals NHS Foundation Trust | Royal Bournemouth Hospital |
| The Royal Oldham Hospital | Pennine Acute Hospitals NHS Trust |
| The Royal Wolverhampton Hospitals NHS Trust | New Cross Hospital |
| The Whittington Hospital NHS Trust | Whittington Hospital |
| Torbay and South Devon Healthcare NHS Foundation Trust | Torbay Hospital |
| United Lincolnshire Hospitals NHS Trust | Grantham and District Hospital |
| United Lincolnshire Hospitals NHS Trust | Lincoln County Hospital |
| United Lincolnshire Hospitals NHS Trust | Pilgrim Hospital |
| University College London Hospitals NHS Foundation Trust | University College Hospital |
| University Hospital of South Manchester NHS Foundation Trust | Wythenshawe Hospital |
| University Hospital of the North Midlands NHS Trust | County Hospital |
| University Hospital of the North Midlands NHS Trust | Royal Stoke University Hospital |
| University Hospital Southampton NHS Foundation Trust | Southampton General Hospital |
| University Hospitals Birmingham NHS Foundation Trust | Queen Elizabeth Hospital Birmingham |
| University Hospitals Bristol NHS Foundation Trust | Bristol Royal Infirmary |
| University Hospitals Coventry and Warwickshire NHS Trust | University Hospital Coventry |
| University Hospitals of Leicester NHS Trust | Glenfield Hospital |
| University Hospitals of Leicester NHS Trust | Leicester Royal Infirmary |
| University Hospitals of Morecambe Bay NHS Foundation Trust | Furness General Hospital |
| University Hospitals of Morecambe Bay NHS Foundation Trust | Royal Lancaster Infirmary |
| Walsall Healthcare NHS Trust | Manor Hospital |
| Warrington and Halton Hospitals NHS Foundation Trust | Warrington Hospital |
| West Hertfordshire Hospitals NHS Trust | Watford General Hospital |

| Trust | Unit |
|--|-------------------------------|
| West Suffolk NHS Foundation Trust | West Suffolk Hospital |
| Western Sussex Hospitals NHS Foundation Trust | St Richard's Hospital |
| Western Sussex Hospitals NHS Foundation Trust | Worthing Hospital |
| Weston Area Health NHS Trust | Weston General Hospital |
| Wirral University Teaching Hospital NHS Foundation Trust | Arrowe Park Hospital |
| Worcestershire Acute Hospitals NHS Trust | Worcestershire Royal Hospital |
| Wrightington, Wigan and Leigh NHS Foundation Trust | Royal Albert Edward Infirmary |
| Wye Valley NHS Trust | County Hospital Hereford |
| Yeovil District Hospital NHS Foundation Trust | Yeovil District Hospital |
| York Teaching Hospital NHS Foundation Trust | Scarborough Hospital |
| York Teaching Hospital NHS Foundation Trust | The York Hospital |

Participating NHS acute units in Wales:

| Trust | Unit |
|--|-------------------------------|
| Abertawe Bro Morgannwg University Health Board | Morrison Hospital |
| Abertawe Bro Morgannwg University Health Board | Princess of Wales Hospital |
| Abertawe Bro Morgannwg University Health Board | Singleton Hospital |
| Aneurin Bevan University Health Board | Nevill Hall Hospital |
| Aneurin Bevan University Health Board | Royal Gwent Hospital |
| Aneurin Bevan University Health Board | Ysbyty Ystrad Fawr Hospital |
| Betsi Cadwaladr University Health Board | Ysbyty Gwynedd |
| Betsi Cadwaladr University Health Board | Glan Clwyd Hospital |
| Cardiff and Vale University Health Board | University Hospital Llandough |
| Cardiff and Vale University Health Board | University Hospital of Wales |
| Cwm Taf University Health Board | Prince Charles Hospital |
| Cwm Taf University Health Board | Royal Glamorgan Hospital |
| Hywel Dda University Health Board | Glangwili General Hospital |
| Hywel Dda University Health Board | Prince Philip Hospital |
| Hywel Dda University Health Board | Withybush General Hospital |

Non-participating NHS acute hospital units

| Trust | Unit |
|---|------------------------------------|
| Barts Health NHS Trust | Whipps Cross University Hospital |
| Betsi Cadwaladr University Health Board | Wrexham Maelor Hospital |
| Cambridge University Hospitals NHS Foundation Trust | Addenbrooke's Hospital |
| Gateshead Health NHS Foundation Trust | Queen Elizabeth Hospital |
| King's College Hospital NHS Foundation Trust | Princess Royal University Hospital |
| Plymouth Hospitals NHS Trust | Derriford Hospital |
| Oxford University Hospitals NHS Foundation Trust | Churchill Hospital |
| Oxford University Hospitals NHS Foundation Trust | Horton General Hospital |
| The Shrewsbury and Telford Hospital NHS Trust | The Princess Royal Hospital |
| The Shrewsbury and Telford Hospital NHS Trust | The Royal Shrewsbury Hospital |

Appendix G: Members of the former secondary care workstream group

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- **Robert A Stone**, National COPD Audit Programme Clinical Lead – Secondary Care Workstream; and Consultant Respiratory Physician, Musgrove Park Hospital, Taunton
- **Noel Baxter**, National COPD Audit Programme Clinical Lead – Primary Care Workstream; and GP Clinical Lead, NHS Southwark CCG
- **Tom Burden**, SpR from Severn region; and Lead for Chest-Reg Network in the South West of England
- **Chris Dyer**, Consultant Geriatrician at the Royal United Hospitals, Bath; and Chair of the British Geriatrics Society Respiratory Special Interest Group
- **Simon Gompertz**, Consultant Respiratory Physician, Queen Elizabeth Hospital Birmingham
- **Ashley Green**, Service Development Manager, British Lung Foundation, London
- **Juliana Holzhauser-Barrie**, National COPD Audit Project Manager, Care Quality Improvement Department, Royal College of Physicians, London
- **John Hurst**, Consultant and Reader, University College London and Royal Free London NHS Foundation Trust
- **Gill Lowrey**, Consultant Respiratory Physician, Royal Derby Hospital
- **Viktoria McMillan**, National COPD Audit Programme Manager, Care Quality Improvement Department, Royal College of Physicians, London
- **Steve Morris**, Professor of Health Economics, University College London
- **Kajal Mortier**, National COPD Audit Project Manager, Care Quality Improvement Department, Royal College of Physicians, London
- **Sandra Olive**, Respiratory Nurse Specialist; and Association of Respiratory Nurses representative
- **Jenni Quint**, Clinical Senior Lecturer in Respiratory Epidemiology, Occupational Medicine and Public Health, National Heart and Lung Institute, Imperial College London; and Honorary Consultant Physician in Respiratory Medicine, Royal Brompton Hospital, London
- **Louise Restrick**, Consultant Respiratory Physician, Whittington Hospital, London
- **James Riordan**, National COPD Audit Programme Coordinator, Quality Improvement Department, Royal College of Physicians, London
- **C Michael Roberts**, National COPD Audit Programme – Programme Clinical Lead; and Clinical Academic Lead for Population Health, UCL Partners
- **Sophie Robinson**, National COPD Audit Programme Coordinator, Care Quality Improvement Department, Royal College of Physicians, London
- **Nick Scriven**, President of the Society for Acute Medicine
- **Laura Searle**, Project Manager, British Thoracic Society, London
- **Michael Steiner**, National COPD Audit Programme Clinical Lead – Pulmonary Rehabilitation Workstream; Honorary Clinical Professor at Loughborough University; and Consultant Respiratory Physician, Glenfield Hospital, Leicester
- **Catherine Thompson**, representative of Association of Chartered Physiotherapists in Respiratory Care (ACPRC); and Head of Patient Experience (Acute Services) at NHS England
- **Sally Welham**, Deputy Chief Executive, British Thoracic Society, London

For further information on the overall audit programme or any of the workstreams, please see our website or contact the national asthma and COPD audit team directly:

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@NACAPaudit
#COPDAudit
#COPDAuditQI

If you would like to join our mailing list and be kept informed of updates and developments in the National Asthma and COPD Audit Programme, please send us your email address and contact details.

Commissioned by:



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Healthcare Quality
Improvement Partnership