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Mouritsen, Jakob Munk; Ehlers, Lars Holger; Kovaleva, Julia; Ahmad, I.; El-Boghdadly, K.

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Review Article

A systematic review and cost effectiveness analysis of reusable vs. single-use flexible bronchoscopes

J. M. Mouritsen, ¹ L. Ehlers, ² J. Kovaleva, ³ I. Ahmad ^{4,5} and K. El-Boghdadly ^{4,5}

- 1 Student, Department of Health Science and Technology, Faculty of Medicine, Aalborg University, Aalborg, Denmark 2 Professor, Danish Center of Healthcare Improvements, Institute of Business and Management, Aalborg University, Aalborg, Denmark
- 3 Clinical Microbiologist/Clinical Pathologist, Sint-Jozefkliniek Bornem and Willebroek, Bornem, Belgium
- 4 Consultant, Guy's and St Thomas' NHS Foundation Trust, London, UK
- 5 Honorary Senior Lecturer, King's College London, UK

Summary

The cost effectiveness of reusable vs. single-use flexible bronchoscopy in the peri-operative setting has yet to be determined. We therefore aimed to determine this and hypothesised that single-use flexible bronchoscopes are cost effective compared with reusable flexible bronchoscopes. We conducted a systematic review of the literature, seeking all reports of cross-contamination or infection following reusable bronchoscope use in any clinical setting. We calculated the incidence of these outcomes and then determined the cost per patient of treating clinical consequences of bronchoscope-induced infection. We also performed a micro-costing analysis to quantify the economics of reusable flexible bronchoscopes in the peri-operative setting from a highthroughput tertiary centre. This produced an accurate estimate of the cost per use of reusable flexible bronchoscopes. We then performed a cost effectiveness analysis, combining the data obtained from the systematic review and micro-costing analysis. We included 16 studies, with a reported incidence of crosscontamination or infection of 2.8%. In the micro-costing analysis, the total cost per use of a reusable flexible bronchoscope was calculated to be £249 sterling. The cost per use of a single-use flexible bronchoscope was £220 sterling. The cost effectiveness analysis demonstrated that reusable flexible bronchoscopes have a cost per patient use of £511 sterling due to the costs of treatment of infection. The findings from this study suggest benefits from the use of single-use flexible bronchoscopes in terms of cost effectiveness, cross-contamination and resource utilisation.

Correspondence to: K. El-Boghdadly Email: elboghdadly@gmail.com Accepted: 19 September 2019

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Twitter: @elboghdadly; @dr_imranahmad

Introduction

Flexible bronchoscopes allow visualisation of the nasopharynx, oropharynx, larynx, trachea and its subsequent divisions for diagnostic and therapeutic purposes. It is estimated that 500,000 bronchoscopic procedures are performed annually in the USA alone [1]. The risk of transmission of infection following bronchoscopy with reusable flexible bronchoscopes is often under-

considered, even when they are reprocessed according to infection control guidelines and recommendations [2–5]. However, the transmission of pathogenic organisms via contaminated reusable flexible bronchoscopes remains an evident risk [6–8], even if appropriate decontamination procedures are adhered to [9]. There are unquantifiable risks of cross-contamination and infection from reusable flexible bronchoscopes, along with uncertainty regarding

their cost effectiveness. Reusable flexible bronchoscopes are often used by anaesthetists to place tracheal tubes, either awake or asleep, and to check adequate positioning of double-lumen tubes. Therefore, cross-contamination risk will also apply to these patients.

Single-use flexible bronchoscopes are delivered sterile and thus should minimise the risk of infection transmission and cross-contamination compared with reusable flexible bronchoscopes. A previously reported cost effectiveness study of single-use flexible bronchoscopes in a typical intensive care unit (ICU) setting in the USA demonstrated that subsequent implementation is cost effective when looking at cross-contamination and potential subsequent infection, and it is associated with increased patient safety [10]. There are several reports of cross-contamination of reusable flexible bronchoscopes due to inappropriate cleaning, disinfection or lack of leak testing and drying [11,12]. These reports do not provide a quantifiable risk for cross-contamination and subsequent infection, but it is accepted that there is a risk and cases are under-reported [8, 13-15]; consequently, the literature lacks a quantified risk of cross-contamination and subsequent infection due to flexible bronchoscopy [8, 16, 17]. Moreover, several microcosting studies of reusable flexible bronchoscopes do not include costs of infections, which is why there is some uncertainty regarding these estimates [18-20].

We therefore aimed to determine the cost per use and cross-contamination risk of reusable flexible bronchoscopes and to ascertain the cost effectiveness of single-use flexible bronchoscopes compared with reusable flexible bronchoscopes in various clinical settings. To achieve this we conducted a micro-costing analysis of flexible bronchoscope utilisation from a high-throughput tertiary centre [21] and a systematic review of the literature. Our primary hypothesis was that single-use flexible bronchoscopes are equally or more cost effective than reusable flexible bronchoscopes.

Methods

The preferred reporting items for systematic reviews and meta-analyses (PRISMA) guidance was adhered to in the conduct of the systematic review [22]. Given an evident risk of patient cross-contamination and infection with reusable flexible bronchoscopes [6,8] and no risk using single-use flexible bronchoscope due to its single-use modality, the effect measure in this cost effectiveness analysis was defined as the avoided risk of infection using single-use compared with reusable flexible bronchoscopes. A comprehensive search strategy was conducted of the PubMed, MEDLINE and Embase databases to identify relevant literature for the risk of

cross-contamination and infection associated with reusable flexible bronchoscopes. The following search terms or combinations were employed: 'bronchoscopy'; 'bronchoscope' determine 'infection' determine 'crossinfection' determine 'pseudo-outbreak' determine 'outbreak' determine 'device contamination' determine and 'hospital infection', including word variations and assorted permutations. English-language studies on humans were included from 1982 to 2018. Retrospective and prospective observational studies were included when studying the risk of cross-contamination and infection post-bronchoscopy. Reference lists of review articles were hand-searched to locate studies that may have been missed in our initial search. Eligibility criteria of studies were met if cross-contamination of patients who underwent bronchoscopic procedures was detected by traditional typing systems based on phenotypes or more recent methods that examine the relatedness of isolates at a molecular level, such as polymerase chain reaction or pulse-field gel electrophoresis [23]. To quantify the risk of infection, we needed the number of: bronchoscopic procedures; patients who underwent bronchoscopy; contaminated patients; and infected patients. Studies were included in the quantitative analysis if at least three of these four variables were reported. Studies were excluded if they had a non-quantifiable risk. A study with a non-quantifiable risk was defined as one in which less than three of the above-mentioned parameters were reported. Other exclusion criteria were other endoscopic procedures (e.g. gastroscopy), if bronchoscope contamination was not detected or reported by recognised typing methods, or if the setting of the studies was not clinically relevant for answering our hypothesis. The risk of infection was used as the measurement of effectiveness. In the event of incomplete data on the number of bronchoscopic procedures and number of patients included, a simple regression method was applied to predict missing data in this large heterogeneous group of patients eligible for bronchoscopy. The relationship was used to predict missing data points within these two variables. Once regression methods were applied, all studies were included for quantitative synthesis. The risk of cross-contamination and infection was determined by a weighted average using a fixed-effects model to reflect a more precise estimate. There are currently no reported cases of cross-contamination using single-use flexible bronchoscopes, which is why the risk is expected to be 0%.

The cost effectiveness of single-use vs. reusable flexible bronchoscopes was estimated by using a literature review to obtain the best available evidence of effects. The effect measure was the risk of infection. The time horizon of the cost-effectiveness analysis was within 1 year. The

micro-costing analysis was conducted at Guy's and St. Thomas' NHS Foundation Trust Department of Anaesthesia. The total cost per use of reusable flexible bronchoscopes for tracheal intubation and double-lumen tube position verification was estimated, and the cost per use of single-use flexible bronchoscopes (including the monitor) was determined. All costs identified were adjusted to pounds sterling (£) in 2017. To determine the present value of capital expenditures, a discount rate of 3.5% was used. Capital acquisition costs were annualised across a 5-year period for bronchoscopes and related equipment, and a 30-year period for buildings.

A decision tree was constructed using Tree Age (2016 version, TreeAge Software, MA, USA), which enabled the comparison of the cost effectiveness of single-use flexible bronchoscopes to reusable flexible bronchoscopes (Fig. 1). The modelling approach was based on principles of good practice for decision analytic modelling in healthcare analyses [24].

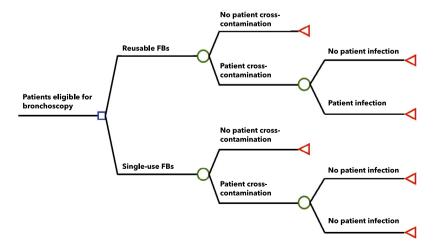
The cost perspective used in this analysis was a UK government third-party payer, and the clinical setting was an anaesthetics department where tracheal intubations and double-lumen tube position verification were frequently carried out. Multiple bronchoscopies for either tracheal intubation or double-lumen tube position verification were examined to determine the resources and costs associated with the procedures. The procedures were monitored in detail from start to finish.

Data obtained retrospectively from various fiscal years (2000–2017) were used to quantify costs for capital acquisitions, repairs, consumables, disposables and service agreements. Additionally, labour time associated with the reprocessing cycles of the reusable flexible bronchoscopes

was meticulously measured. The department has 19 reusable flexible bronchoscopes. Of these, 12 are used for tracheal intubation (either awake or asleep), whereas seven are reserved for double-lumen tube position verification. The activities of performing bronchoscopy are dispersed across various operating theatres on one floor, whereas the reprocessing is divided on two different floors. The reusable flexible bronchoscopes undergo a precleaning cycle on the same floor as the operating theatre using equipment including detergents and brushes, followed by a second-stage manual clean performed in a central cleaning facility on another floor. Costs were estimated using the mean annual number of bronchoscopic procedures and reprocessing volumes done by the automated endoscope reprocessors in this tertiary hospital.

In the Supporting Information (Data S1), all collected cost data from the micro-costing analysis are presented in detail. This includes: (1) capital and repair costs of reusable flexible bronchoscopes and rack systems; (2) capital and repair costs of reprocessing capital and additional equipment; (3) time measurements of the specific reprocessing steps; (4) average cost of labour-related reprocessing; (5) reprocessing equipment and costs incurred; and (6) the allocation keys and reason for usage [25].

The costs of the clinical outcome were determined by identifying treatment costs related to the clinical manifestations of a post-bronchoscopic contamination or infection. The incidence of postoperative pulmonary complications (including pneumonia and sepsis) is estimated to be up to 23% in an unselected group of patients having general anaesthesia [26]. This figure is higher in patients undergoing thoracic or head and neck surgery [27], which are the cohort of patients in whom



 $\textbf{Figure 1} \quad \text{Decision tree model used in this cost effectiveness analysis}.$

flexible bronchoscopes are most commonly used. However, there are no data directly demonstrating bronchoscopeinduced cross-contamination or infection in these patients, as infection is often assumed to be multifactorial. We were therefore only able to use published data of infection and cross-contamination from the ICU or elective endoscopy or bronchoscopy population. The cost of the clinical outcome was estimated from the studies that were included according to the clinical manifestations that were reported in included studies, such as respiratory tract infection prophylaxis and therapy [28-36] and sepsis [33, 35]. In the cohort of patients of interest for this study, we considered ventilator-associated pneumonia, sepsis and communityacquired pneumonia as suitable outcomes. The average cost of ventilator-associated pneumonia was identified from a review of 28 US community hospitals to be £25,426 sterling per patient [37]. From a systematic review of hospital-related cost of sepsis, the treatment-related costs were identified as £27,123 sterling per patient [38]. The costs of inpatient and outpatient community-acquired pneumonia were estimated from > 28,000 communityacquired pneumonia episodes from a large US database study at £13,151 and £1948, respectively, per patient [39]. The weighted average was defined as the treatment-related costs per patient infected. This value was imputed in the cost-effectiveness analysis. A summary of all costs and effect inputs for the cost-effectiveness analysis is presented in Table 1.

Using the results from the literature review and the micro-costing analysis to compute the cost-effectiveness analysis, a base-case result was generated. Sensitivity analyses were undertaken to capture uncertainty within parameters and to provide sufficient insight to decision-

makers. Uncertainty is captured through deterministic and probabilistic sensitivity analyses. One-way (univariate) sensitivity analyses were applied to all parameters in the model to test its robustness by examining the impact on the incremental cost effectiveness ratio. All cost parameters were varied by \pm 50%. Considering the average period of an 8-month investigation across the 16 studies, the effect parameters were varied from a low value of 0% risk to a high value of 20% risk. A scenario analysis was conducted from a previous Delphi approach to the general risk of patient cross-contamination and infection [10]. Furthermore, the impact of the incremental cost-effectiveness ratio was observed by varying the amortisation period of capital investments related to bronchoscopy materials to 10 years from the previous 5 years.

The probabilistic sensitivity analysis quantifies the overall uncertainty within parameters using pre-specified distributions (Table 1). The modality of the probabilistic sensitivity analysis was a second-order Monte Carlo simulation with 10,000 iterations of the mean incremental cost-effectiveness ratios. These 10,000 iterations were drawn up in a cost effectiveness scatterplot to represent the expected avoidance of infection risk using a single-use compared with reusable flexible bonchoscopes.

According to International Guidelines in Health Economics, the mean is used as it is the only relevant measure for economic decision making [40]. In economic calculations, we aim to capture the uncertainty of the sample mean, that is, parameter uncertainty, rather than variability or heterogeneity, that is, stochastic variability. The uncertainty in the expected mean is the standard error (SE) [40]. Consequently, we report all cost data as mean (SE).

Table 1 Inputs for the cost effectiveness model. Base-case value, the standard error (SE) and the distribution are provided.

Parameters	Base-case value (SE)	Distribution
Effects		
Risk of patient contamination using a reusable FB	0.153 (0.009)	Beta
Risk of subsequent infection using a reusable FB	0.181 (0.018)	Beta
Risk of patient contamination using a single-use FB	0.0 (0.001)	Beta
Risk of subsequent infection using a single-use FB	0.0 (0.001)	Beta
Costs		
$Capital\ cost\ per\ use\ of\ a\ reusable\ FB\ (reusable\ FB, stack\ systems, reprocessing\ capital)$	£116.4a (29.10)	Gamma
Repair cost per use of a reusable FB (reusable FB, stack systems, reprocessing capital)	£92.9*(23.20)	Gamma
Reprocessing cost per use a reusable FB (labour time and equipment)	£39.9*(10.00)	Gamma
Cost of per clinical outcome [37–39] (per patient infection)	£9,454(£1,158)	Gamma
Cost per use of a single-use FB, including monitor	£220 (21.80)	Gamma

^aSE was not estimated. However, a conservative approach was taken by varying the parameter with a SE of 25%. FB, flexible bronchoscope.

Results

We identified 890 citations, of which 12 were duplicates. Seven additional studies were considered through hand-searching of two review articles [7, 8]. Across numerous studies a non-quantifiable risk was identified [41–62], thus not fulfilling the inclusion criteria for the quantitative analysis. After screening based on title and abstract and full-text review, we identified 16 studies for quantitative analysis of the cross-contamination and infection risk (Fig. 2) [12, 28–36, 63–68]. The setting of these studies were patients who underwent bronchoscopy in a hospital intensive care or respiratory unit setting, or during bronchoscopy or endoscopy in the UK, USA, France, Spain,

Australia or Taiwan (Table 2). The study designs were prospective observational and retrospective studies, and the period of contamination and infection investigation of patients undertaken was from one to 23 months [30, 33]. A total of 2351 patients underwent 3120 various bronchoscopic procedures. Of these procedures, 476 cases of patient contamination were detected. Eighty-six of these patients were reported to have a bronchoscope-linked manifestation of infection, including pneumonia or other respiratory tract infection. Seven of the included studies contained a missing data-point relating to the number of bronchoscopic procedures (five studies) and the number of patients who underwent a bronchoscopy (two studies)

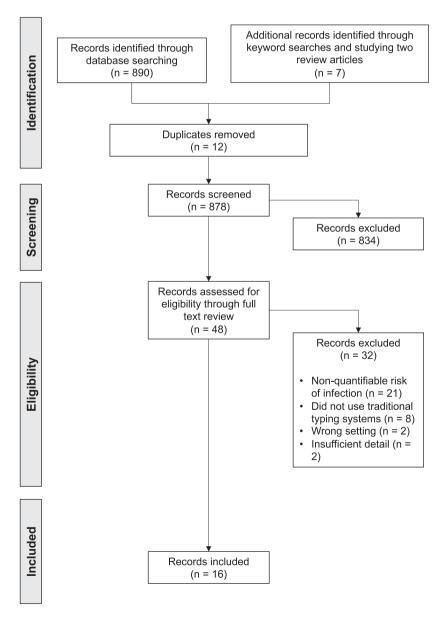


Figure 2 Study flow chart.

 Table 2
 Sixteen studies identified for quantitative analysis of the cross-contamination and infection risk. Values are number or mean (SD).

		-	,									
Source	Study design	Period of investigation (months)	Country	Setting	Patient age (years)	Proce- dures	Patients	Cases of contami- nation	Cases of infection	Detection of contami- nation	Typing system used	Infection(s)
Blanc et al. [68]	Observational	9	USA	Y Y	٩	410	299a	35	0	Yes	Ribotyping	Noinfections
Botana-Rial et al. [30]	Observational	-	Spain	BU	∢ Z	154	118a	39	21	≺es	REP-PCR	Treatment to prevent the development of pneumonia
Chroneou et al. [63]	Observational	2	NSA	₹ Z	71 (49–86)	76a	57	6	0	Yes	REP-PCR	Noinfections
DiazGranados et al. [35]	Observational	2	USA	BU	₹ Z	27a	20	12	2	Yes	PFGE	Pneumonia and sepsis
Waite et al. [64]	Retrospective	12	¥	ICN	NA	63a	47	18	0	Yes	PFGE	Noinfections
Wang et al.[31]	Retrospective	4	Taiwan	₹ Z	60 (45–79)	163a	123	18	80	≺es	AFB	Patients were treated as mycobacterial infected
Silva et al. [12]	Retrospective	22	NSA	EU	NA A	429a	324	41	0	Yes	Ribotyping	Noinfections
Nye et al. [65]	Retrospective	9	¥	3	61 (40–80)	28	7	7	0	Yes	Culturing and lipid analysis	Noinfections
Guy et al. [32]	Retrospective	7	France	ICO	62 (49–73)	216	157	10	8	≺es	PFGE	Treatment to prevent the development of pneumonia
Sammartino et al. [36]	Retrospective	m	NSA	AN A	56 (36–76)	19	19	11	_	Yes	Serotyping	Pneumonia
Campagnaro et al. [66]	Retrospective	2	Australia	Y Y	ΑN	92	92	12	0	Yes	DNA probing	Noinfections
Corne et al. [28]	Retrospective	∞	France	ICU	NA A	61	36	16	4	Yes	PFGE	Pneumonia
Kirschke et al. [34]	Retrospective	4	NSA	AN A	59 (24–88)	99	09	20	_	Yes	PFGE	Pneumonia
Srinivasan et al. [33]	Retrospective	23	USA	E	∢ Z	992	414	39	39	≺es ≺	PFGE	Pneumonias, sepsis, respiratory tract infection
Pappas et al. [29]	Retrospective	11	NSA	A A	ΑN	195	187	72	2	Yes	Culturing	No data
Cêtre et al. [67]	Observational/ retrospective	∞	France	E	₹ Z	453	418	117	0	Yes	PFGE	Noinfections
Total						3120	2351	476	98			
						i						

AFB, acid-fast bacillus testing; BU, bronchoscopic unit; DNA, deoxyribonucleic acid; EU, endoscopic unit; ICU, intensive care unit; NA, not available; PFGE, pulsed-field gel electrophoresis; REP-PCR, repetitive extragenic palindromic-polymerase chain reaction.

Predicted data-point

[12, 30, 31, 35, 63, 64, 68]. These missing data-points were predicted using a linear regression method. There was a strong correlation between the number of patients who underwent bronchoscopy and the number of bronchoscopic procedures (r = 0.92) (see Supporting Information, Data S2). From this, 118 and 299 patients were predicted, respectively, for two of the studies with a missing data point [30, 68]. A strong correlation was also demonstrated between the number of bronchoscopic procedures carried out and the number of patients who underwent bronchoscopy (r = 0.93). Five studies contained a missing data-point in terms of number of bronchoscopic procedures, which were therefore calculated as 163, 429, 27, 76 and 63, respectively [12, 31, 35, 63, 64].

From the included studies, the differentiation between a pre-existing infection and a flexible bronchoscope-related infection was determined using traditional bacterial recognition or more recent methods that examine the association of isolates at a molecular level (Table 2).

One thousand annual bronchoscopic procedures are performed by the Department of Anaesthesia at Guy's Hospital. The reprocessing volumes by the automated endoscope reprocessor were 10,075 cycles per year. The results from the micro-costing analysis revealed a mean (SE) capital cost per use of a reusable flexible bronchoscope at £116.40 (£29.10), whereas the repair and reprocessing cost per use of a reusable flexible bronchoscope was estimated at £92.90 (£23.20) and £39.90 (£10.00), respectively. This equates to a total cost per use of a reusable flexible bronchoscope at £249.20 sterling. The mean (SE) cost per use of a single-use flexible bronchoscope were provided by Ambu (Ambu® aScope™ 4, Copenhagen, Denmark) at £220.00 sterling (£21.80), including the monitor.

In the Supporting Information (Data S1), a detailed overview and description are provided of all costs incurred, and the allocation keys that were employed to more accurately reflect reality.

In the cost-effectiveness analysis, we found reusable flexible bronchoscopes to have a mean (SE) cost per patient of £511.00 sterling (£59.60), with an associated risk of infection of 2.8%. The mean (SE) cost per patient with single-use flexible bronchoscopes are estimated at £220.00 (£21.80) and a 0% risk of infection. Base-case results indicate a net saving of £291.00 to hospitals and an avoided risk of infection of patients undergoing bronchoscopy at 2.8% with single-use flexible bronchoscopes compared with reusable flexible bronchoscopes. The base-case incremental cost-effectiveness ratio is -£10,505, which is interpreted as the cost of avoided patient infection.

Varying cost inputs by \pm 50% did not have a significant impact on the expected value of the incremental cost-effectiveness ratio, which varied from $-\pounds15,232$ to $-\pounds5,778$. From Fig. 3, the cost parameter with the greatest impact on cost effectiveness is the cost of clinical outcome, whereas the parameter with the lowest impact is the reprocessing cost per use of a reusable flexible bronchoscope.

Considering the average of 8 months of investigation across the 16 studies of patient contamination and infection, one-way sensitivity results from the CEA and a scenario analysis is presented in Table 3.

The probabilistic sensitivity analysis was indicative of a potential net savings to hospitals ranging from £34 to 577 sterling per use and eliminating the risk of infection of approximately 1.71–4.07% using single-use flexible bronchoscopes compared with reusable flexible bronchoscopes (Fig. 4).

Discussion

This systematic review and cost-effectiveness analysis found that single-use flexible bronchoscopes are cost effective and associated with a lower risk of infection compared with reusable flexible bronchoscopes. Sensitivity analyses support these findings. Our systematic review demonstrated that the risk of patient infection post-bronchoscopy was 2.8%, with a cost per use of a reusable flexible bronchoscope of £249 sterling and of a single-use flexible bronchoscope of £220 sterling. Our cost-effectiveness analysis demonstrated that reusable flexible bronchoscopes have a cost per patient use of £511 sterling due to the potential costs of treatment of infection.

The risk for patient contamination (15%) and infection (18%) resulted in a 2.8% risk of patient infection post-bronchoscopy. In a previous study, a Delphi approach was used to estimate the general risk of patient contamination (3.4%) and infection (21%) in critical care settings [10]. This generated a risk of post-bronchoscopy infection of 0.7%. The estimate from this present study (2.8%) is of high accuracy because patient contamination and infection were linked to bronchoscopes and the data were sourced from international, multicentre settings with more than 2300 patients undergoing approximately 3100 various procedures.

The cost per use of reusable flexible bronchoscopes for tracheal intubation in a UK hospital carrying out 141 flexible bronchoscope-assisted tracheal intubations annually was previously estimated as £340 sterling [69]. In this present study, the total cost per use of reusable flexible bronchoscopes in an institution performing 1000 annual

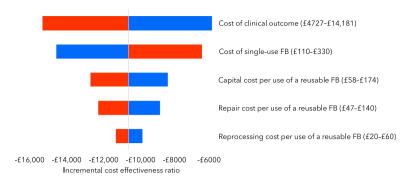


Figure 3 Tornado chart showing multiple one-way (univariate) sensitivity analyses of cost input parameters varied by \pm 50%. The incremental cost-effectiveness ratio (ICER) midpoint is -£10,505 sterling and is equal to the base-case result from the cost-effectiveness analysis. Low values (blue) for cost of clinical outcome, capital, repair and reprocessing cost per use of a reusable flexible bronchoscope increase the ICER, whereas high values (red) reduce the ICER. Low values (blue) for cost of a single-use bronchoscope reduce the ICER, whereas high values (red) increase the ICER. FB, flexible bronchoscope.

 Table 3
 Base-case result and one-way sensitivity analyses of effect parameters.

Description	Difference in cost	Difference in effects	ICER (cost per avoided patient infection)	
Base-case	-£291	2.8%	Dominant	
One-way sensitivity of effects				
Cross-contamination risk of 0%	-£29	No difference	Dominant	
Cross-contamination risk of 20%	-£371	3.6%	Dominant	
Infection risk of 0%	-£29	No difference	Dominant	
Infection risk of 20%	-£318	3.1%	Dominant	
One-way sensitivity of amortisation of capital investments				
Capital investments amortised across 10 years	-£239	2.8%	Dominant	
Scenario analysis using estimates of cross-contamination and infection risk obtained from a Delphi approach[10]				
Cross-contamination risk of 3.38% and infection risk of 21.3%	-£97	0.7%	Dominant	

ICER, incremental cost effectiveness ratio.

procedures was estimated to be £249 sterling. The cost per use for capital, reprocessing and repairs are highly dependent on local and clinical setting. The reprocessing costs are dependent on the length of non-usage before reusable flexible bronchoscopes need reprocessing again. This is a variable time, but can be as low as 12 h [5]. Staffing costs and use of productive working hours are also factors that may have an economic impact. Furthermore, repair costs per use of reusable flexible bronchoscopes are also highly dependent on the local setting. McCahon and Whynes conducted an analysis at a teaching hospital with repair cost per use of £146 sterling [69], which contrasts to the repair cost per use at Guy's Hospital of £93 sterling. This could be due to service agreements to cover repairs of all capital equipment in some institutions but not others. Finally, capital costs per use are dependent on economies of scale advantages, that is, the volume of annual

bronchoscopies. Comparing McCahon and Whynes data to that in Guy's Hospital, the capital cost per use was £141 and £114, respectively. Other costing analyses have been conducted with total cost per use ranging from £111 to £540 sterling [18–20]. From these studies, the clinical setting in terms of capital and repair expenditures play an important role when determining the capital and repair cost per use, whereas the reprocessing costs were similar between settings.

There are two main strengths of this study, the first being the micro-costing analysis as the cost comparison between reusable and single-use equipment is complex. Numerous overhead cost elements must be considered, and this study captures more than previous studies [18–20]. The other main strength is the fact that this is the first study to identify risk of patient contamination and infection from the published literature.

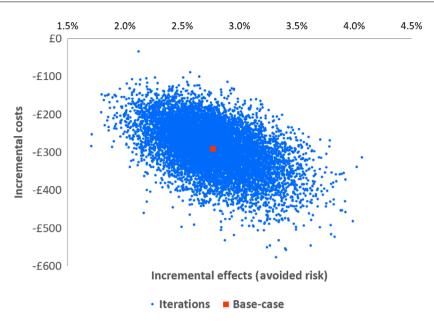


Figure 4 Probabilistic sensitivity analysis with 10,000 iterations (blue) and the base-case value (red).

In general, cost analyses lack precision in terms of including all relevant overhead costs [18-20]. This is difficult to manage because numerous elements contribute to the overall cost per use estimate, such as water consumption, electricity, training of new personnel in cleaning techniques, maintaining updated and compliant guidelines on reprocessing, handling of automated endoscope reprocessor cycle failures, among others. Moreover, the cost of the tracking systems is often left out [18-20]. If all relevant indirect and overhead costs were identified and included in those analyses, evidently it would add to the cost per use. To advance the precision of the cost per use estimate in this present micro-costing analysis, some of the overhead costs mentioned above could have been included. If done so, it would have added to the cost effectiveness of single-use flexible bronchoscopes. Findings from sensitivity analyses of cost per use of singleuse flexible bronchoscopes and reusable flexible bronchoscopes support the cost effectiveness of single-use flexible bronchoscopes, even when varying cost parameters \pm 50%.

In the institution examined in this study, the availability of reusable flexible bronchoscopes remains limited due to the unplanned requirement for flexible bronchoscopes, the additional time required for reprocessing and the ongoing requirement for bronchoscopes to be repaired. Availability was still an issue, thus investment in a subset of single-use flexible bronchoscope for emergencies was undertaken. The availability of single-use flexible bronchoscopes is constant as long as stocks are replenished, but the

availability of reusable flexible bronchoscopes are hampered by the need for reprocessing.

The environmental impact of clinical care is also an element to consider. The disposable equipment, chemical detergents, water, electricity, and other resources used during the reprocessing cycles have an environmental impact, whereas single-use flexible bronchoscopes are disposed of after each use. From research comparing carbon dioxide (CO₂) emissions and resource consumption from a single-use flexible bronchoscope (Ambu aScope[™] 4) to a reusable flexible bronchoscope, results show that the materials used for reprocessing are substantial when comparing the two types of bronchoscopes [70]. The use of cleaning materials and personal protective equipment contributes to a comparable or potentially higher material and energy consumption as well as emissions of CO2 equivalents and value of resource consumption for reusable compared with single-use flexible bronchoscopes [70].

This study has several limitations. Our cost-effectiveness analysis show that, ceteris paribus, the single-use technology is superior in terms of costs and patient safety, and a mix of single-use and reusable equipment may be the only realistic alternative. But not all overhead costs related to reusable equipment will be possible to eliminate, and the predicted savings may be smaller under such circumstances. Further research should be conducted to investigate the cost effectiveness of a mixed usage strategy for single-use and reusable bronchoscopes. Another potential limitation is that we were unable to perform a formal risk of bias assessment of the included studies

as no appropriate, validated tool was suitable. An additional limitation was that we included data from patients presenting for outpatient bronchoscopy, who may sometimes have systemic pathology, could be immunocompromised and are, therefore, prone to infection [6-8]. However, the cohort of patients sought in the microcosting analysis also included patients undergoing thoracic surgical procedures and patients undergoing major head and neck surgery, both of which may be associated with high risk of infection transmission and immunocompromise. Moreover, the risk of transmission of infection remains, regardless of patient baseline characteristics. Furthermore, our cost calculations are primarily relevant to the perioperative setting, but the published data used come from outpatient bronchoscopic and critical care settings, and therefore there may be some discrepancy that is unaccounted for due to this assumption. There was heterogeneity in the lengths of investigations (1-23 months), patient contamination rates (4.6-58%) and infection risk (0-100%) among included studies. The variation was partly accounted for by utilising a fixed-effects model. When varying risks of patient cross-contamination and infection in the sensitivity analysis, single-use flexible bronchoscopes remained financially superior to reusable flexible bronchoscopes. In addition, one of the assumptions that were made was that the risk of infection of single-use flexible bronchoscopes was 0%. However, there always remain unaccounted-for risks of infection transmission, even with single-use flexible bronchoscopes, due to unsterile operators and surrounding equipment or re-use in the same patient [71]. Finally, there remains a previously unreported risk of nosocomial infection from tracheal intubation in elective patients who are not critically unwell and who are intubated for a limited duration of time, which might account for some of the incidence of infection after bronchoscopy. However, given that there are virtually no data reporting nosocomial infection in these settings, this may have a limited impact on our data. Moreover, the data we used on the published incidence of infection could be secondary to tracheal intubation rather than the use of a flexible bronchoscope. However, given that only 44 of the included patients with evidence of infection were in critical care settings, it is unlikely that the incidence of infection could be primarily attributed to non-bronchoscopic sources.

In conclusion, our systematic review has demonstrated that the risk of patient infection following bronchoscopy with reusable flexible bronchoscopes is significant, warranting a need for guidelines on reprocessing to be stricter to ensure greater patient safety [8, 9, 72]. The total

cost per use of a reusable flexible bronchoscope was calculated to be £249. The cost per use of a single-use flexible bronchoscope was £220 sterling. When considering the risk of infection in the cost analysis, reusable flexible bronchoscopes have a mean cost per patient of £511 sterling and an associated risk of infection at 2.8%. The findings from this study suggest benefits of single-use flexible bronchoscopes in terms of cost effectiveness, crosscontamination and resource utilisation.

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Supporting Information

Additional supporting information may be found online via the journal website.

Data \$1. Micro-costing analysis.

Data S2. Data prediction.