

Prognostic study to detect ailments, including cancer, during solid organ transplantation to prevent postoperative complications using perioperative antibiotics therapy

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ABSTRACT

Objective: The purpose of this research was to examine the use of preoperative antibiotic therapy in solid organ transplantation to prevent postoperative complications.

Methods: The data examined through PsycINFO and CINAHL conference papers, searches for resources that would be useful for our study. A conceptual study of randomized controlled trials and quasi-RCTs on antibiotic prophylaxis for solid organ transplant patients to avoid post-operative complication at any time after transplantation. Antibiotic resistance and surgical site infections were major findings. A model with unpredictable effects was employed to compute the RR and 95% CIs. Key findings were antibiotic resistance and surgical site infections. Using a random-effects model, we calculated RRs and 95% CIs.

Results: 617 randomized participants. 237 randomized individuals compared antibiotics to no antibiotics, while 370 compared extended-duration to short-duration antibiotics. The evidence is extremely weak that antibiotics minimize surgical site infections. The other results were quite uncertain. There is some evidence that individuals who have had solid organ transplants may benefit from using extended-duration antibiotics to avoid surgical site infections. None of the 7 studies examined possible harmful effects on grafts, heart disease, malignancy, life expectancy, haematological and biological indicators, length of hospital stay, cost of assistance, or hospitalisation charges.

Conclusion: Available information on the effectiveness of pre-operative antibiotic prophylaxis for organ transplantation is of very low-quality owing to methodological constraints, bias risk, and high heterogeneity. More excellent RCTs with sufficient power would improve clinical practice.

Keywords: perioperative antibiotic, postoperative complications, solid organ transplantation, malignancy, Randomized Controlled Trials (RCTs), quasi-RCTs

INTRODUCTION

The most common cause of morbidity and mortality after solid organ donation is infection. Depending on the donated organ and the period, postoperative wound infections have been documented to happen in anywhere between 3% and 53% of transplant recipients. An infection at the surgical site develops after surgery in the area of the body where the operation was performed. This might involve the treating wound's erythema and returning to the operating room for wound debridement [1]. The surgery site infection rates for small bowel transplants, which may reach over 90% when the prosthetic mesh is utilized, are the highest, followed by liver, stomach, and kidney transplantation. Infections at the surgical site are linked to significant morbidity and have been shown to increase readmission rates, increase hospital costs by over 100%, and lengthen the typical hospital stay by seven days. Furthermore, solid organ transplant patients who suffer surgical site infections have higher odds of graft failure and mortality. Also, those who get solid organ transplants are at significant risk of getting infections from bacteria that are resistant to antibiotics. Multidrug-resistant infections have been linked to higher morbidity and mortality, especially in solid organ transplant patients. Thus, methods must be created to reduce surgical site infections after transplantation. The treatments and research reported need to be divided into different groups depending on the kind of patient (organ), the period (era), and the analysis of microbiological data [2]. Initially, this prophylactic prescription was known as "antibiotic prophylaxis," but more recently, the term "preventive antibiotic treatment" had been established to denote exclusively the preventive use of antibiotics in healthy individuals to prevent early failure and the emergence of postoperative infections [3]. In the past, patients received implant treatment that were completely dentures, and then it was expanded to include patients who were just partly dentures. However, the quantity of bone available for placing dental implants is often decreased due to the loss of the alveolar ridges in the maxilla unless a reconstructive phase is carried out, and there are several types of bone atrophy and associated treatment protocols. Maximizing surgical rates and minimizing problems undergoing lift procedures often necessitates an interdisciplinary strategy including many experts in the pre-surgical period [4]. Before maxillary sinus elevation, certain anatomical abnormalities and pathological disorders, such as inflammation infective processes symptoms of cancer-related illnesses, should be addressed. Postoperative infections are uncommon, occurring

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between 2% to 5.6% of the time, and there is no differentiation between real sinus infections and infections from sinus grafts [5].

Research evaluated the efficacy of various antibiotic prophylaxis regimens in comparison to a placebo in terms of potential postoperative problems resulting from the surgical extraction of impacted lower third molars [6]. Their findings demonstrated that using antibiotics to prevent postoperative problems was a good idea and that using them just after the operation has been performed has no advantages over using them both before and after. Study offered an update on the current landscape of the use of antimicrobial stewardship methods for improving PP [7]. A study assessed SOT as a possible contributor to complications after THA for ONFH [8]. The potentially crippling disorder ONFH often necessitates complete THA replacement. Postoperative problems after THA for osteoarthritis were more likely to occur in patients who have had SOT. In the highlight new research on the use of extracellular vesicles as biomarkers for various illnesses and talk about how they could be used in clinical practice [9].

Preclinical research found that reduced inflammation and oxidative stress led to better graft survival [10]. Results from human research are still early, and some have not succeeded in transferring the positive outcomes from animal studies to the clinical context. Also, there was a lot of variation across studies in terms of the method of administration, how the donor or recipient was treated, if graft flushing was used, and whether cadaveric grafts were used. The introduction of current pharmacotherapeutic strategies, including those integrated into apps and features of electronic medical records, was given by study linked to reduced readmission rates decreased hospital length of stay, decreased frequency and severity of postoperative complications, and cost or revenue impacts [11]. Results from tests might provide suggestions for tailoring medication for each patient in light of their particular genetic makeup. In study, they assemble the most recent data supporting the use of micro biome research in LT, with an emphasis on infections and biliary problems brought on by multidrug-resistant microorganisms [12]. Results of these investigations have shown that in individuals with biliary problems after liver transplantation, harmful bacteria may be found in specimens of bile or bile ducts. The purpose of this research was to examine the use of preoperative antibiotic therapy in solid organ transplantation to prevent postoperative complications.

The remainder of this research is structured in the following manner:

- Part 2 introduces the method of the material
- The result analyses are in Part 3
- Part 4 contains the discussion
- Part 5 contains the conclusion
- Part 6 contains the limitation and future scope

MATERIALS AND METHODS

All RCTs and quasi-RCTs examine the safety, effectiveness, or both of perioperative treatment of antibiotic-related postoperative complications in Patients Receiving SOT.

Inclusion criteria

Patients of any age receive a solid-organ transplant, including

those with graft loss from several recipients (e.g. kidney-pancreas).

Exclusion criteria

Post-natal care women.

Interventions of antibiotic medication

Any research using an antibiotic was considered. Inquiries were conducted into the following pairs of comparisons.

- The use of any antibiotic drug *vs.* placebo or no therapy
- Every antibiotic *vs.* every other antibiotic
- Utilizing a single antibiotic at low and high doses
- Treatment with short- or long-term antibiotics
- Comparison of the effects of antibiotics given by mouth with those given intravenously

Data sources

Studies in the register are indexed from the following databases.

- CINAHL is a comprehensive database that indexes literature related to nursing and allied health professions.
- The database primarily focuses on literature relevant to healthcare practitioners, educators, and researchers in nursing and allied health fields. It includes articles on clinical practices, patient care, healthcare administration, and evidence-based medicine.
- PsycINFO is a leading database for literature in psychology and related fields. The database includes journal articles, books, dissertations, conference papers, and technical reports.

Cochrane Kidney and Transplant's scope is used to identify studies for the Register via searches of CINAHL and PsycINFO. The search tactics used and the publications, conferences, and alerts that were manually searched may all be found on the cochrane kidney and transplant website.

Search strategy

Titles and abstracts of articles pertinent to the assessment were obtained using the search approach specified. 4 writers separately revised the titles and abstracts, excluding any studies that didn't apply but keeping those that seemed like they would have useful data or information. To determine whether articles met the inclusion criteria, 4 writers independently revised all abstracts and, if required, the entire texts of the publications that were retrieved. Where 4 authors could not agree on whether or not an article should be included, the deciding vote was taken by a research assistant.

Data extraction

Each contributor extracted data on their own using industry-standard templates. Research published in periodicals other than English has to be translated before analysis. In cases where more than one report on the same subject had been published, this study pooled the reports and selected the one with the most comprehensive data for our analysis. If results were only available in pre-publication drafts, we utilized them. Instances, where there was a disparity between the several print runs, were marked.

RESULTS

Study process

After looking through the specialized registry, we found 13

separate reports. The first screening led to the inclusion of 7 studies (13 reports) and the exclusion of 2 studies (2 reports). There is currently just one active research. Figure 1 shows the overall flow for the study.

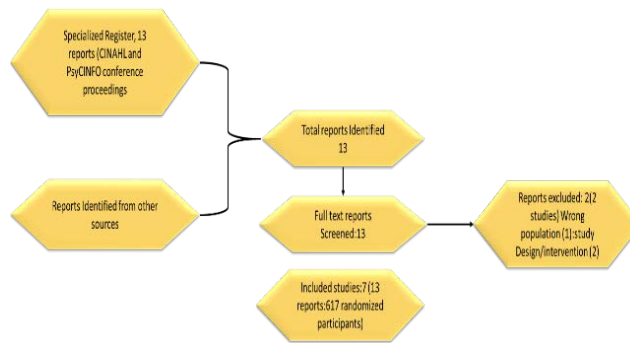


Fig. 1. Overall flow of the study

The conceptual comprised 7 trials with a total of 617 randomized participants. In three studies, 237 patients with previous SOT were given antibiotics for the prevention of postoperative complications, and the results were compared to those who received a placebo or no antibiotics. 3 trials with a total of 370 randomized participants compared the efficacy of longer courses of antibiotics

to those of shorter courses in reducing the risk of infection at surgical sites for people who had had solid-organ transplants.

Antibiotics for reducing the risk of postoperative complications in SOT patients are shown in figure 2.

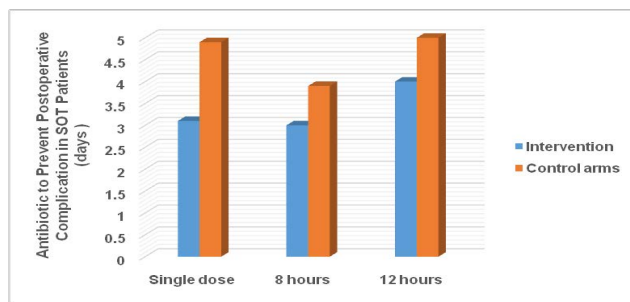


Fig. 2. Antibiotics to prevent post-operation complications SOT patients

Cefazolin or cefotaxime every 12 hours for 3 days to 5 days vs. a single dose, piperacillin/tazobactam every eight hours for three days vs. a single dose, and this medication with doxycycline as opposed to cefazolin and sulbactam. It wasn't specified the length of either therapy arms may endure. Patients who had liver resection were disqualified since the focus of the research should have been on liver transplantation; patients who presented with a fever 30 days after receiving a kidney transplant should have been evaluated for the effectiveness of antimicrobial treatment, etc.

the incidence of surgical site infection is quite weak, however. Due to the poor quality of the evidence, it is not known whether antibiotics lower the overall mortality rate or not. The confidence of the data from single research is quite low, it is not known if antibiotics reduce graft loss, even death with a functioning transplant. Antibiotics may or may not lower UTI incidence; this is unclear due to the poor quality of the available data. It is not known if antibiotics lessen the occurrence of septicemia since the evidence is rated as extremely poor in confidence. As the data from single research has been rated as extremely poor in confidence, we do not know whether antibiotics reduce the occurrence of pneumonia or not.

To compare the effectiveness of antibiotics against no therapy for avoiding infection at the surgical site in recipients of solid organ transplants, see table 1. The data suggesting that antibiotics lower

| Tab. 1. Antibiotic without any antibiotic for SOT patients' postoperative complication | Findings | Comparative Impact (95%CI) | Confidence in a Record | Anticipated Immediate Effect (95% CI) | |
|--|--|----------------------------|------------------------|---------------------------------------|---------------------------|
| | | | | The Risk of no Antibiotics | Risk with Antibiotics |
| | Graft loss | RR2.92 (0.13 to 67.97) | Extremely minimal | 41 per 1,000 | 0 per 1,000 |
| | Surgical site infections* Time frame: 2 weeks to 6 weeks | RR0.46 (0.24 to 0.9) | Extremely minimal | 316 per 1,000 | 138 per 1,000 (66 to 268) |
| | Death (any case) Time frame: 2 weeks to 6 weeks | RR0.30 (0.06 to 3.73) | Extremely minimal | 134 per 1,000 | 41 per 1,000 (3 to 473) |
| | Other infection: pneumonia Timeframe: up to 6 weeks | RR0.54 (0.24 to 1.36) | Extremely minimal | 475 per 1,000 | 253 per 1,000 (99 to 621) |

| | | | | |
|---|-----------------------|-------------------|------------------|----------------------------|
| Adverse reactions | - | - | Never recognized | Never recognized |
| Other infection: septicaemia Time frame: 2 weeks to 6 weeks | RR0.51 (0.14 to 2.23) | Extremely minimal | 96 per 1,000 | 51 per 1,000 (10 to 204) |
| Other infection: UTI Time frame: 2 weeks to 6 weeks | RR0.90 (0.72 to 1.18) | Extremely minimal | 624 per 1,000 | 549 per 1,000 (421 to 706) |

Antibiotics for avoiding infections at the transplant recipient's surgical site are included in table 2. There is not enough data to conclude that prolonged use of antibiotics helps prevent infections at surgical sites. Due to the limited confidence in the findings of a single research, we do not know whether longer-lasting antibiotics lower the risk of surgical site infections in patients who received just a kidney in a transplant. The data from single research is quite weak, so we don't know whether administering extended-duration antibiotics to people who've had liver transplants but no other organs are effective in lowering the rate of surgical-site infections. LT recipients who were given piperacillin/tazobactam for 3 days were more likely to have VRE establish intra-abdominally, a liver relocate patient was more likely to develop intra-abdominal and blood vancomycin-resistant enterococci after receiving an im-

mediate dose of piperacillin/tazobactam. Whether the presence of resistant organisms is a result of antibiotic treatment before transplantation or if these individuals naturally harbored VRE makes it difficult to determine. The data from a single trial is relatively weak, so we don't know whether using antibiotics for longer periods improves mortality. Due to the lack of evidence, we cannot tell whether prolonged antibiotics minimize graft loss, such as death, in functional transplants. The data from a single trial is quite weak, so we don't know whether using antibiotics for a longer period lowers UTIs. The data suggesting that using antibiotics for longer periods lowers the occurrence of septicemia is quite weak, however. As the data from single research has been rated as extremely poor quality, we do not know whether using antibiotics for longer periods helps reduce pneumonia.

Tab. 2. Prolonged vs. short-term antibiotics for postoperative complications in SOT patients

| Findings | Comparative Impact (95%CI) | Confidence in a Record | Anticipated Immediate Effect (95%CI) | |
|---|----------------------------|------------------------|---|--|
| | | | The risk with Extended-Duration Antibiotics | The Risk with Short-Duration Antibiotics |
| Surgical site infections: kidney transplant recipients | RR0.52 (0.05 to 5.48) | Extremely minimal | 14 per 1,000 (1to108) | 23 per 1,000 |
| Surgical site infections (All) Time frame: 30 days | RR1.21 (0.59 to 2.50) | Extremely minimal | 91 per 1,000 (43 to 183) | 77 per 1,000 |
| Inadequate graft survival Estimated duration: 30 days | RR 0.97 (0.51 to 2.03) | Extremely minimal | 93 per 1,000 (46 to 187) | 96 per 1,000 |
| Death (any cause) Time frame: 30 days | RR0.23 (0.03 to 3.97) | Extremely minimal | 13 per 1,000 (0 to 168) | 49per 1,000 |
| Surgical site infections: liver transplant recipients Estimated duration: 30 days | RR1.33 (0.63 to 2.84) | Extremely minimal | 249per 1,000 (114 to 527) | 193per 1,000 |
| Another complication: septicaemia Estimated duration: 30 days | RR 0.92(0.53 to 1.61) | Extremely minimal | 136 per 1,000 (74 to 230) | 149 per 1,000 |
| Another complication: UTI Estimated duration: 30 days | RR 0.51 (0.14 to 1.88) | Extremely minimal | 66 per 1,000 (16 to 231) | 129 per 1,000 |
| Adverse reactions | - | - | Never recognized | Never recognized |
| Another complication: pneumonia Estimated duration: 30 days | RR 11.69 (0.61 TO 189.79) | Extremely minimal | 5/50** | No events |

Transplants occur on average once every year, as shown in figure 3. Figure 3 show that most participating facilities conducted more than 10 solid organ transplantations, with a few being high-ur-

gency transplantations and a range in the number of live-related SOT. In the operating room, both pediatric and general critical care units provided immediate postoperative treatment.

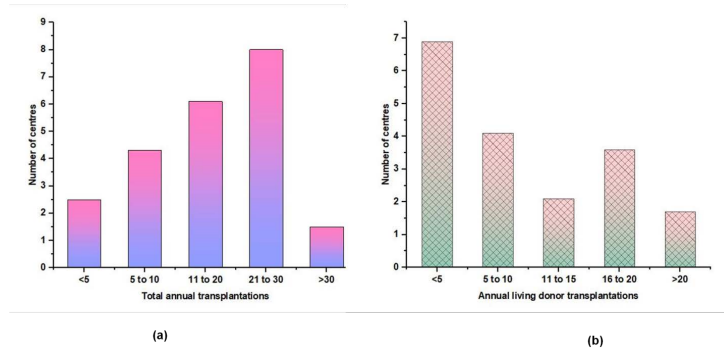


Fig. 3. Yearly transplantations in cooperating hospitals

Figure 4 depicts the perioperative antibiotic prophylaxis of risk factors. Antibiotic prophylaxis before surgery in children who have had a liver transplant varies in length depending on their risk factors. Treatment-related variables (e.g., hospital length of stay, antibiotic use, or the use of an intra-abdominal patch) are considered risk factors. Antibiotic therapy was often administered for a shorter or longer amount of time than that was considered normal by certain centers, depending on the patient's unique risk level. Around 14 locations provided extensive data on potential dangers. These included MDR colonization, the presence of an

abdominal patch, the progression of c-reactive protein and procalcitonin levels after surgery, antimicrobial therapy before the transplant, the presence of preexisting, indwelling central lines, the presence of ascites after surgery, the patient's age, the length of hospitalization before transplantation, and the number of previous surgical procedures. Centers with fewer transplants per year (≤ 20) and centers with more transplant patients per year (>20) exhibited comparable durations of prophylaxis when divided by a yearly number of transplants. However, the former utilized a narrow range antibiotic more often.

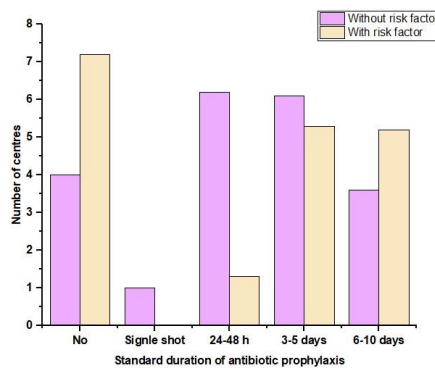


Fig. 4. Perioperative antibiotic prophylaxis of risk factors

DISCUSSION

The results of this systematic analysis indicate that there is inadequate high-quality information to draw a firm conclusion about whether or not antibiotics decrease the frequency of infections at the surgical site. The 7 studies that were included had highly heterogeneous results, had a high risk of bias, and had poor methodological designs. In particular, there were many separate studies, conflicting similarities with variable outcome measures, and little information on problems like Overuse of antibiotics [13]. Without a single element of the primary studies published any of the SONG initiative's main objectives. We were unable to investigate and accurately quantify possible problems emerging from perioperative antibiotic usage due to the limits of the available information. In general, it is challenging to translate the findings of this Cochrane study into clinical practice. First off, since kidney and liver transplant patients were the focus of the included RCTs, the findings may not generalize to other SOT, which can carry different risks of perioperative infection. Patients who have had multi-visceral transplants may be more at risk for infections and may need longer-lasting preoperative medications. Second, five of the studies had their first publication date before the year 2000. Given the radical shifts that have occurred in recipient and donor selection, immune modulator regimens, and microbial resistance

patterns and pathogenesis, it is possible that it no longer applies to contemporary transplant treatment. None of the seven concepts evaluated donor infection while limiting antibiotics. Positive donor meconium-stained amniotic cultures were connected to longer critical care unit stays and worse post-transplant survival rates in lung transplant recipients. Fourth, only three of the eight papers that were included in the analysis described the microorganisms' antimicrobial treatment resistance patterns. As has been shown in recipients of heart, lung, liver, and kidney transplants, colonization may increase the likelihood of surgical site infections. The eight studies' use of eight different, non-standard criteria of infections at the surgical site raises the possibility of categorization bias [14]. Actively physicians looking into surgical site infections have a big impact on whether they are found. It is challenging to diagnose surgical site infections in the hospital environment since they are often found after departure and managed again in the community [15]. Due to suboptimal methodological planning, notably the omission of mechanisms for randomization and allocation concealment, the quality of the available evidence was compromised. A significant percentage of trials also had a substantial risk of efficiency and detection biases. Studies were often constrained by a tiny sample and an uncertain follow-up period, which diminished the review's strength [16]. These limitations imply that further research is likely to alter our confidence in effect estimates.

CONCLUSION

This research strength is the rigorous CINAHL, and PsycINFO search that was conducted to locate only RCTs and quasi-RCTs that met the evaluations pre-determined inclusion criteria. Since no participants had a financial stake in the outcome of this research, we cannot rule out the potential that an underlying conceptual conflict affected the researchers' assessment of the information and, by extension, their conclusions. There are parts of other studies that we agree with and parts that we disagree with. While antibiotics have been shown to reduce surgical site infections, this investigation demonstrates that the evidence is limited. While there are broad guidelines for the prophylactic use of antibiotics in the perioperative period preceding heart, lung, liver, kidney, and kidney-pancreas transplantation, no such guidelines have been documented for SOT patients.

LIMITATION AND FUTURE SCOPE

Antibiotics are no longer routinely given to SOT patients to prevent post-operative complications since the data supporting this practice is so weak. There has to be more research done on the topic of long-term vs. short-term antibiotic treatment.

Further research comparing long-acting antibiotics to their shorter-acting counterparts for the prevention of surgical site infections would be useful. The current randomized investigation has the potential to provide light on the present unknowns. Nevertheless, a variety of organism-specific transplants should be taken into account so that more accurate estimations may be used to guide treatment. Research in the future should use relevant samples (such as rectal swabs and urine specimens) and a systematic approach to determine not only the baseline degree of antimicrobial resistance but also the change in drug resistance antibiotic treatment.

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