

World Journal of *Experimental Medicine*

Quarterly Volume 14 Number 4 December 20, 2024



EDITORIAL

Cheng CH, Hao WR, Cheng TH. Harnessing aryl hydrocarbon receptor dynamics: Unveiling therapeutic pathways in esophageal squamous cell carcinoma. *World J Exp Med* 2024; 14(4): 98599 [DOI: [10.5493/wjem.v14.i4.98599](https://doi.org/10.5493/wjem.v14.i4.98599)]

REVIEW

Wibowo DP, Agustiningsih A, Jayanti S, Sukowati CHC, El Khobar KE. Exploring the impact of hepatitis B immunoglobulin and antiviral interventions to reduce vertical transmission of hepatitis B virus. *World J Exp Med* 2024; 14(4): 95960 [DOI: [10.5493/wjem.v14.i4.95960](https://doi.org/10.5493/wjem.v14.i4.95960)]

Maiti A, Mondal S, Choudhury S, Bandopadhyay A, Mukherjee S, Sikdar N. Oncometabolites in pancreatic cancer: Strategies and its implications. *World J Exp Med* 2024; 14(4): 96005 [DOI: [10.5493/wjem.v14.i4.96005](https://doi.org/10.5493/wjem.v14.i4.96005)]

Suleman A, Aluyi-Osa G, Ashipa F, Spadea L, Gagliano C, D'Esposito F, Zeppieri M, Musa M. Autologous blood in the management of ocular surface disorders. *World J Exp Med* 2024; 14(4): 96412 [DOI: [10.5493/wjem.v14.i4.96412](https://doi.org/10.5493/wjem.v14.i4.96412)]

Cavaillon JM, Chaudry IH. Facing stress and inflammation: From the cell to the planet. *World J Exp Med* 2024; 14(4): 96422 [DOI: [10.5493/wjem.v14.i4.96422](https://doi.org/10.5493/wjem.v14.i4.96422)]

MINIREVIEWS

de Paulo CB, Miglino MA, Castelucci P. Perspectives on the extracellular matrix in inflammatory bowel disease and bowel decellularization protocols. *World J Exp Med* 2024; 14(4): 97179 [DOI: [10.5493/wjem.v14.i4.97179](https://doi.org/10.5493/wjem.v14.i4.97179)]

Schuch LF, Silveira FM, Pereira-Prado V, Sicco E, Pandiar D, Villarroel-Dorrego M, Bologna-Molina R. Clinicopathological and molecular insights into odontogenic tumors associated with syndromes: A comprehensive review. *World J Exp Med* 2024; 14(4): 98005 [DOI: [10.5493/wjem.v14.i4.98005](https://doi.org/10.5493/wjem.v14.i4.98005)]

Arora A, Morya AK, Gupta PC, Menia NK, Nishant P, Gupta V. Intravitreal therapy for the management of diabetic retinopathy: A concise review. *World J Exp Med* 2024; 14(4): 99235 [DOI: [10.5493/wjem.v14.i4.99235](https://doi.org/10.5493/wjem.v14.i4.99235)]

Sridhar GR, Gumpeny L. Melanocortin 4 receptor mutation in obesity. *World J Exp Med* 2024; 14(4): 99239 [DOI: [10.5493/wjem.v14.i4.99239](https://doi.org/10.5493/wjem.v14.i4.99239)]

ORIGINAL ARTICLE**Retrospective Study**

Salzillo C, Basile R, Cazzato G, Ingravallo G, Marzullo A. Value of autopsy in the modern age: Discrepancy between clinical and autopsy diagnoses. *World J Exp Med* 2024; 14(4): 95147 [DOI: [10.5493/wjem.v14.i4.95147](https://doi.org/10.5493/wjem.v14.i4.95147)]

Alshaikhsalama A, Archer H, Xi Y, Ljuhar R, Wells JE, Chhabra A. HIPPO artificial intelligence: Correlating automated radiographic femoroacetabular measurements with patient-reported outcomes in developmental hip dysplasia. *World J Exp Med* 2024; 14(4): 99359 [DOI: [10.5493/wjem.v14.i4.99359](https://doi.org/10.5493/wjem.v14.i4.99359)]

Clinical Trials Study

Seif El-Din Z, Afify M, Zayed E, Elsabaawy D, Tharwa ES, Elsharawy A, Abdelsameea E, Rady MA. Dapagliflozin as an oral antihyperglycemic agent in the management of diabetes mellitus in patients with liver cirrhosis. *World J Exp Med* 2024; 14(4): 95272 [DOI: [10.5493/wjem.v14.i4.95272](https://doi.org/10.5493/wjem.v14.i4.95272)]

META-ANALYSIS

Tarar ZI, Farooq U, Inayat F, Basida SD, Ibrahim F, Gandhi M, Nawaz G, Afzal A, Chaudhary AJ, Kamal F, Ali AH, Ghouri YA. Statins decrease the risk of hepatocellular carcinoma in metabolic dysfunction-associated steatotic liver disease: A systematic review and meta-analysis. *World J Exp Med* 2024; 14(4): 98543 [DOI: [10.5493/wjem.v14.i4.98543](https://doi.org/10.5493/wjem.v14.i4.98543)]

LETTER TO THE EDITOR

Bangolo AI, Wadhwani N. Comprehensive analysis of the impact of primary percutaneous coronary intervention on patients with ST-segment elevation myocardial infarction. *World J Exp Med* 2024; 14(4): 94845 [DOI: [10.5493/wjem.v14.i4.94845](https://doi.org/10.5493/wjem.v14.i4.94845)]

Xiang Z, Li JR, Wan WM, Li SH, Wu J. Familial hypercholesterolemia: Current limitations and future breakthroughs. *World J Exp Med* 2024; 14(4): 99968 [DOI: [10.5493/wjem.v14.i4.99968](https://doi.org/10.5493/wjem.v14.i4.99968)]

ABOUT COVER

Peer Reviewer of *World Journal of Experimental Medicine*, Ramachandra Barik, Professor, Department of Cardiology, All India Institute of Medical Sciences, Bhubaneswar 751019, India. cardioramachandra@gmail.com

AIMS AND SCOPE

The primary aim of the *World Journal of Experimental Medicine (WJEM, World J Exp Med)* is to provide scholars and readers from various fields of experimental medicine with a platform to publish high-quality basic and clinical research articles and communicate their research findings online.

WJEM mainly publishes articles reporting research results and findings obtained in the field of experimental medicine and covering a wide range of topics including clinical laboratory medicine (applied and basic research in hematology, body fluid examination, cytomorphology, genetic diagnosis of hematological disorders, thrombosis and hemostasis, and blood typing and transfusion), biochemical examination (applied and basic research in laboratory automation and information system, biochemical methodology, and biochemical diagnostics), etc.

INDEXING/ABSTRACTING

The *WJEM* is now abstracted and indexed in PubMed, PubMed Central, Scopus, Reference Citation Analysis, China Science and Technology Journal Database, and Superstar Journals Database. The *WJEM's* CiteScore for 2023 is 1.7 and Scopus CiteScore rank 2023: Internal medicine is 109/167.

RESPONSIBLE EDITORS FOR THIS ISSUE

Production Editor: *Lai Zhang*, Production Department Director: *Xu Guo*, Cover Editor: *Ji-Hong Liu*.

NAME OF JOURNAL

World Journal of Experimental Medicine

ISSN

ISSN 2220-315x (online)

LAUNCH DATE

December 20, 2011

FREQUENCY

Quarterly

EDITORS-IN-CHIEF

Jian Wu

EXECUTIVE ASSOCIATE EDITORS-IN-CHIEF

Fang Gong

EDITORIAL BOARD MEMBERS

<https://www.wjgnet.com/2220-315x/editorialboard.htm>

PUBLICATION DATE

December 20, 2024

COPYRIGHT

© 2024 Baishideng Publishing Group Inc

PUBLISHING PARTNER

Department of Clinical Laboratory, The Affiliated Suzhou Hospital of Nanjing Medical University, Suzhou Municipal Hospital

INSTRUCTIONS TO AUTHORS

<https://www.wjgnet.com/bpg/gerinfo/204>

GUIDELINES FOR ETHICS DOCUMENTS

<https://www.wjgnet.com/bpg/GerInfo/287>

GUIDELINES FOR NON-NATIVE SPEAKERS OF ENGLISH

<https://www.wjgnet.com/bpg/gerinfo/240>

PUBLICATION ETHICS

<https://www.wjgnet.com/bpg/GerInfo/288>

PUBLICATION MISCONDUCT

<https://www.wjgnet.com/bpg/gerinfo/208>

POLICY OF CO-AUTHORS

<https://www.wjgnet.com/bpg/GerInfo/310>

ARTICLE PROCESSING CHARGE

<https://www.wjgnet.com/bpg/gerinfo/242>

STEPS FOR SUBMITTING MANUSCRIPTS

<https://www.wjgnet.com/bpg/GerInfo/239>

ONLINE SUBMISSION

<https://www.f6publishing.com>

PUBLISHING PARTNER'S OFFICIAL WEBSITE

<http://www.smh.cc/home2020/page/index/index.html>

Retrospective Study

Value of autopsy in the modern age: Discrepancy between clinical and autopsy diagnoses

Cecilia Salzillo, Roberta Basile, Gerardo Cazzato, Giuseppe Ingravallo, Andrea Marzullo

Specialty type: Pathology**Provenance and peer review:**

Invited article; Externally peer reviewed.

Peer-review model: Single blind**Peer-review report's classification****Scientific Quality:** Grade B, Grade B**Novelty:** Grade B, Grade B**Creativity or Innovation:** Grade C, Grade C**Scientific Significance:** Grade B, Grade B**P-Reviewer:** Zhou ZB**Received:** April 2, 2024**Revised:** July 19, 2024**Accepted:** August 5, 2024**Published online:** December 20, 2024**Processing time:** 211 Days and 18.2 Hours**Cecilia Salzillo, Gerardo Cazzato, Giuseppe Ingravallo, Andrea Marzullo**, Department of Precision and Regenerative Medicine and Ionian Area, Pathology Unit, University of Bari "Aldo Moro", Bari 70121, Italy**Cecilia Salzillo**, Department of Experimental Medicine, PhD Course in Public Health, University of Campania "Luigi Vanvitelli", Naples 80138, Italy**Roberta Basile**, Course in Biomedical Laboratory Technique, University of Bari "Aldo Moro", Bari 70121, Italy**Co-first authors:** Cecilia Salzillo and Roberta Basile.**Corresponding author:** Cecilia Salzillo, MD, MScCVP, Expert in Pathology, Department of Precision and Regenerative Medicine and Ionian Area, Pathology Unit, University of Bari "Aldo Moro", Piazza Giulio Cesare 11, Bari 70121, Italy. c.salzillo@studenti.uniba.it**Abstract****BACKGROUND**

Autopsy is a medical procedure that consists of the examination of the corpse to determine the cause of death and obtain information on pathological conditions or injuries. In recent years, there has been a reduction in hospital autopsies and an increase in forensic autopsies.

AIM

To evaluate the utility of autopsy in the modern age and the discrepancy between clinical and autopsy diagnoses.

METHODS

A retrospective observational study was conducted on the reports of all 645 hospital autopsies performed at Polyclinic of Bari from 2006 to 2021.

RESULTS

Group A, 2006-2009, 174 cases were studied: 58% male, 58% adults, 55% neonatology; pulmonary disease was the cause of death in 23% of cases; and there was a discrepancy between clinical and autopsy diagnosis in 55% of cases. Group B, 2010-2013, 119 cases: 52% male, 46% infants, 48% neonatology; pulmonary disease was the cause of death in 25% of cases; and there was a discrepancy between clinical and autopsy diagnosis in 56% of cases. Group C, 2014-2017, 168

cases: sex equality, 37% infants, 25% gynecology; pulmonary disease was the cause of death in 24% of cases; and there was a discrepancy between clinical and autopsy diagnosis in 58% of cases. Group D, 2018-2021, 184 cases: 56% male, 38% adult, 32% gynecology; pulmonary disease was the cause of death in 27% of cases; and there was a discrepancy between clinical and autopsy diagnosis in 58% of cases.

CONCLUSION

The study of hospital autopsies reveals a 56.75% discrepancy between clinical diagnosis and autopsy, highlighting the importance of autopsies, especially for fetal and neonatal diseases, which represent 59% of cases.

Key Words: Hospital autopsy; Modern age; Clinical diagnosis; Autopsy diagnosis; Public health

©The Author(s) 2024. Published by Baishideng Publishing Group Inc. All rights reserved.

Core Tip: The hospital autopsy is useful in the modern age, especially for the diagnosis of fetal and neonatal pathologies. Genetic and non-genetic diagnoses are important to future parents for subsequent pregnancies and can thus be studied.

Citation: Salzillo C, Basile R, Cazzato G, Ingravallo G, Marzullo A. Value of autopsy in the modern age: Discrepancy between clinical and autopsy diagnoses. *World J Exp Med* 2024; 14(4): 95147

URL: <https://www.wjgnet.com/2220-315x/full/v14/i4/95147.htm>

DOI: <https://dx.doi.org/10.5493/wjem.v14.i4.95147>

INTRODUCTION

The term "autopsy" is derived from Greek and means "to see for oneself". There are two types of autopsy: hospital autopsy and forensic autopsy.

The first law authorizing human dissection was enacted in 1231 by Frederick II. In the 16th century, Andreas Vesalius initiated the modern study of anatomy with the publication of "De Humani Corporis Fabrica"[1]. Giovanni Morgagni correlated clinical symptoms with organic changes, introducing the anatomical-clinical concept, and published "De sedibus et causis morborum per anatomen indagatis"[2].

In the 19th century, Karl von Rokitansky performed over 30000 autopsies and wrote the "Handbook of Pathological Anatomy"[3], while Rudolf Virchow, the founder of modern pathology, introduced detailed autopsy techniques[4]. Maurice Letulle described the technique of mass organ removal[5], and Albrecht Ghon introduced the technique of *en bloc* removal.

Over the last decades, there has been a significant decrease in the number of hospital autopsies and an increase in judicial autopsies[6]. This phenomenon may be due to various factors, including changes in medical practices, economic issues and the evolution of diagnostic techniques. Hospital autopsies are used as a tool to improve the quality of care and diagnostic accuracy. However, their utilization has declined due to budget constraints and growing confidence in imaging technologies[7,8]. Forensic autopsies are performed to determine the cause of death for forensic purposes. Their frequency has increased due to a greater emphasis on medico-legal responsibility and the need for thorough investigations in cases of suspicious or violent death[9,10]. This underlines how our society has an interest in the legal aspect rather than in knowing the cause of death.

In this study, we aimed to analyze 645 hospital autopsy cases from 2006 to 2021 retrieved from the digital archive of the Pathology Unit, Department of Precision and Regenerative Medicine and Ionian Area, Polyclinic of Bari, to study the rate of concordance between clinical and autopsy diagnosis and evaluating whether the execution of hospital autopsies is helpful in the modern age.

MATERIALS AND METHODS

Study design

In our retrospective observational study, we analyzed the autopsy case history of the Pathology Unit, Department of Precision and Regenerative Medicine and Ionian Area, Polyclinic of Bari.

Patient data source

We used the digitalized archive of autopsy reports between 2006 and 2021 for a total of 645 cases, and all cases in the archive were included in the study.

Subgrouping for analysis

The 645 cases were divided into groups of 4 years to make the samples uniform and comparable: group A: 2006-2009 of 174 cases, group B: 2010-2013 of 119 cases, group C: 2014-2017 of 168 cases, and group D: 2018-2021 of 184 cases.

Each subgroups was divided by age: Fetus < 180 days or < 6 months gestational age, infant/newborn 1 year of age, child/adolescent 1-16 years of age, and adult > 16 years of age.

The total divisions into subgroups included the following: Sex, age, specialty, autopsy diagnosis, and correlation between clinical and autopsy diagnosis.

To analyze the pathological diagnosis, we grouped the cause of death into: Cardiovascular, infectious, miscellaneous, neoplastic, placental, pulmonary, and syndromes/malformations.

Finally, we analyzed the discrepancy between clinical and autopsy diagnoses.

RESULTS

Group A

Out of 174 cases in group A (Table 1), 58% were male, 41% female, and 1% undefined, of which 58% were adults, 1% children, 29% infants, and 12% fetuses. Out of these 174, the specialty were: neonatology 55%, gynecology 19%, internal medicine 10%, external 7%, and other departments 9%. The cause of death was classified as pulmonary disease in 23%, syndromes and/or malformations 10%, infections 8%, cardiovascular diseases 6%, placental disease 4%, neoplasms 4%, and miscellaneous 45%. In 55% of the analyzed cases, there was discrepancy between clinical and autopsy diagnoses (Figure 1A).

Group B

Group B (Table 2) comprised 119 cases; 51% male, 48% female, and 1% undefined, of which 31% were adults, 6% children, 46% infants, and 17% fetuses. The specialty of the origin of the deceased was 48% neonatology, 19% gynecology, 10% cardiac surgery, 12% general surgery, 4% neurology, and 7% others. The cause of death was categorized as 25% pulmonary diseases, 15% cardiovascular diseases, 11% syndromes and/or malformations, 9% infectious diseases, 3% neoplasms, 1% placental pathology, and 36% miscellaneous. In 56% of the cases, a discrepancy between clinical and autopsy diagnoses was observed (Figure 1B).

Group C

Group C (Table 3) comprised 168 cases, of which 50% were male and 50% female; 26% were adults, 1% were children, 37% infants, and 36% fetuses. The origin of the deceased was 30% gynecology, 25% neonatology, 17% cardiac surgery, 6% emergency room, 5% general surgery, 4% from other regional hospitals, and 13% others. The cause of death was classified as 24% pulmonary diseases, 15% cardiovascular diseases, 11% syndromes and/or malformations, 6% placental pathology, 3% infections, 2% neoplasms, and 39% miscellaneous. In 58% of the cases, there was a discrepancy between clinical and autopsy diagnoses (Figure 1C).

Group D

Group D (Table 4) comprised 184 cases; 56% male, 43% female, and 1% undefined; 38% were adults, 3% children, 26% infants, and 33% fetuses. Of these 184 cases, the specialty of origin was 32% gynecology, 21% other regional hospitals, 21% neonatology, 14% emergency department, and 12% others. In 27% of patients, the cause of death involved pulmonary diseases, 23% cardiovascular diseases, 6% syndromes and/or malformations, 5% placental disease, 4% infections, 2% neoplasms, and 33% miscellaneous. In 58% of the cases, there was a discrepancy between clinical and pathological diagnoses (Figure 1D).

Statistical analysis of changes over time

From our study it is evident that sex distribution has remained constant over time. In fact, the percentage of males showed only slight variations, going from 58% in group A to 56% in group D; whereas the percentage of females remained stable, fluctuating between 41% and 50%. This stability suggests that there were no significant changes in the sex distribution over the study period.

We observed an important change in age distribution. The percentage of newborns and fetuses increased over time, with a peak in Group C where newborns were 37% of cases and fetuses were 36% of cases; however, the percentage of adults significantly decreased, from 58% in Group A to 38% in Group D. This trend demonstrates the growing attention of autopsies to pediatric and neonatal cases in recent years.

Neonatology is one of the main specialties involved in autopsies, but the percentage fell from 55% in group A to 21% in group D. In contrast, gynecology had a significant increase, becoming the predominant specialty in group D with 32%. This change reflects possible changes in clinical practices and autopsy patient populations.

Lung and cardiovascular diseases consistently have high rates as causes of death across all groups. However, diagnoses classified as "miscellaneous" made up a significant proportion in each group, with a slight decrease from 45% in Group A to 33% in Group D. This broad and generic category of diagnoses could include a variety of conditions that do not fit into major disease categories, likely reflecting the complexity and diversity of autopsy cases examined.

The percentage of discrepancy between the clinical diagnosis and the autopsy diagnosis remained high and constant throughout the study period, with a variation between 55% and 58%. This discrepancy highlights the importance of

Table 1 Group A: Analyzing the 174 hospital autopsies, 2006-2009

Group A	Sex	Age	Specialty	Autopsy diagnosis	Discrepancy
174 cases, 2006-2009	58% male, 41% female, 1% undefined	58% adults, 29% infants, 12% fetuses, 1% children	55% neonatology, 19% gynecology, 10% internal medicine, 9% others, 7% external	23% pulmonary, 10% syndromes/malformations, 8% infections, 6% cardiovascular, 4% placental, 4% neoplasms, 45% miscellaneous	55%

Table 2 Group B: Analyzing the 119 hospital autopsies, 2010-2013

Group B	Sex	Age	Specialty	Autopsy diagnosis	Discrepancy
119 cases, 2010-2013	51% male, 48% female, 1% undefined	31% adults, 46% infants, 17% fetuses, 6% children	48% neonatology, 19% gynecology, 12% general surgery, 10% cardiac surgery, 7% others, 4% neurology	25% pulmonary, 15% cardiovascular, 11% syndromes/malformations, 9% infectious, 3% neoplasms, 1% placental, 36% miscellaneous	56%

Table 3 Group C: Analyzing the 168 hospital autopsies, 2014-2017

Group C	Sex	Age	Specialty	Autopsy diagnosis	Discrepancy
168 cases, 2014-2017	50% male, 50% female	37% infants, 36% fetuses, 26% adults, 1% children	30% gynecology, 25% neonatology, 17% cardiac surgery, 13% others, 6% emergency room, 5% general surgery, 4% external	24% pulmonary, 15% cardiovascular, 11% syndromes/malformations, 6% placental, 3% infections, 2% neoplasms, 39% miscellaneous	58%

Table 4 Group D: Analyzing the 184 hospital autopsies, 2018-2021

Group D	Sex	Age	Specialty	Autopsy diagnosis	Discrepancy
184 cases, 2018-2021	56% male, 43% female, 1% undefined	38% adults, 33% fetuses, 26% infants, 3% children	32% gynecology, 21% neonatology, 21% external, 14% emergency room, 12% others	27% pulmonary, 23% cardiovascular, 6% syndromes/malformations, 5% placental, 4% infections, 2% neoplasms, 33% miscellaneous	58%

autopsies as a crucial diagnostic tool, underscoring the limitations of clinical diagnoses and the utility of autopsy in revealing conditions not identified during life.

DISCUSSION

By analyzing the 645 hospital autopsies (Table 5), we observed a predominance of 53.75% males, 59% newborns and fetuses. The most frequent specialty of origin was 37.25% neonatology and 25% gynecology. The cause of death in our study was primarily pulmonary diseases followed by cardiovascular diseases, and the discrepancy between clinical and autopsy diagnoses was over 56.75%.

Our study highlights that hospital autopsies were mainly performed in fetuses and infants, and the requirement of autopsy in adults was progressively reduced.

The decrease in hospital autopsies in adult patients is probably due to the excessive trust in medical diagnostic technology, whereby it is believed that the autopsy does not provide any additional information that is not already known at the time of death[11].

Although some studies have underlined the importance of carrying out a hospital autopsy[6,11,12], medical doctors often do not recognize the importance and therefore do not explain the advantages to the deceased's relatives. Autopsy is a fundamental tool for understanding pathological processes, the effectiveness of treatments, the correct diagnostic approaches, and for preventing medical errors and supporting public health[13,14].

In a study conducted in Sweden by Friberg *et al*[15] on 2410 hospital autopsies of adult patients, there was an overall reduction in the request for autopsy examination with prevalence of cardiovascular disease as the cause of death, and with a discrepancy of more than 30% between clinical diagnosis and autopsy. The authors highlighted how the hospital autopsy provides information about the disease and cause of death that is likely unknown to the doctor and presumably to the relatives of the deceased and explained how this can have a negative impact on public health[15,16].

In a prospective cohort study, Latten *et al*[17] investigated the relationship between clinical cause of death, a history of cancer, and the rate of medical autopsies. The authors observed that the autopsy rate was positively correlated with the number of causes of death, suggesting a higher value of interest in autopsies in complex medical cases. According to the authors, healthcare and individuals would benefit from an increase in post-mortem investigations[17].

Table 5 Analyzing the 645 hospital autopsies, 2006-2021

Group	Sex	Age	Specialty	Autopsy diagnosis	Discrepancy
A + B + C + D = 645 cases, 2006-2021	53.75% male, 45.5% female, 0.75% undefined	38.25% adults, 34.5% infants, 24.5% fetuses, 2.75% children	37.25% neonatology, 25% gynecology, 10.25% others, 8% external, 6.75% cardiac surgery, 5% emergency room, 4.25% general surgery, 2.5% internal medicine, 1% neurology	24.75% pulmonary diseases, 14.75% cardiovascular diseases, 9.5% syndromes/malformations, 6% infections, 4% placental disease, 2.75% neoplasms, 38.25% miscellaneous	56.75%

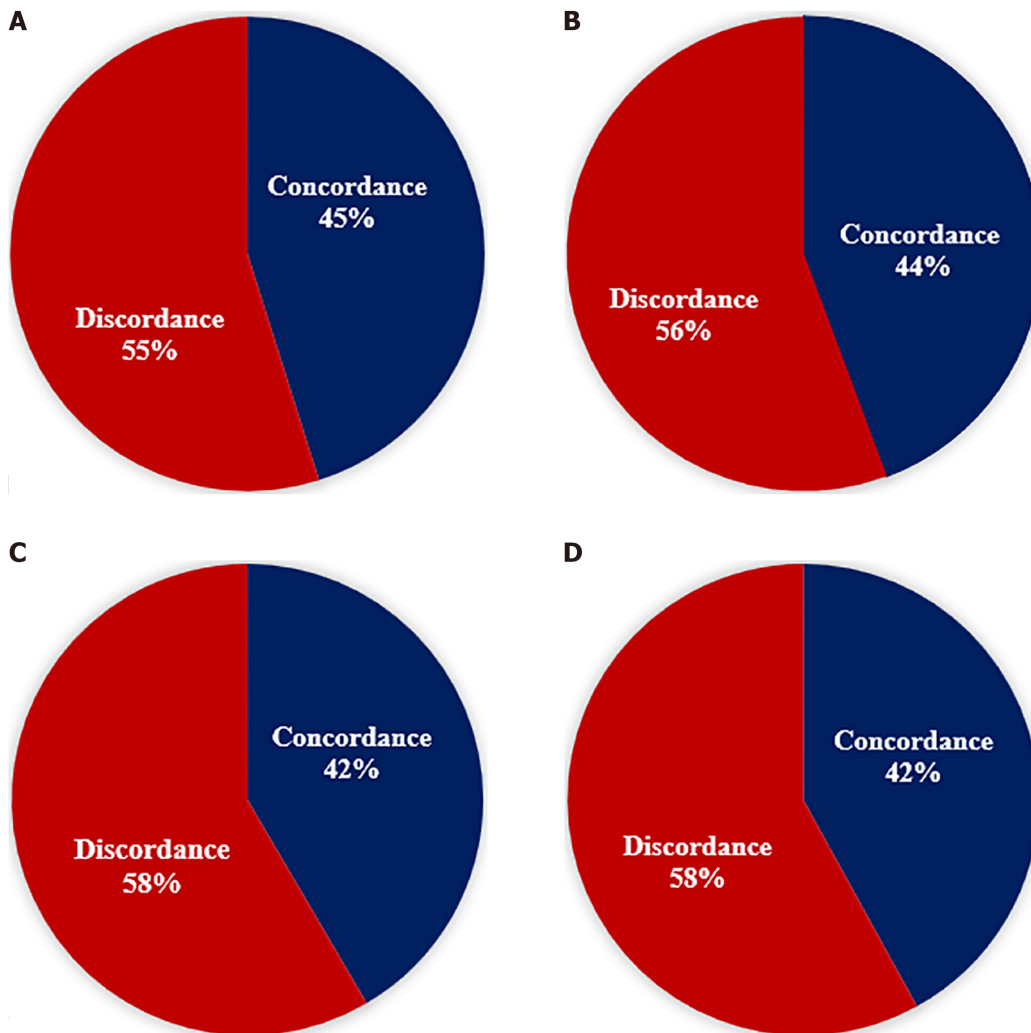


Figure 1 Discrepancy between clinical and autopsy diagnosis. A: Group A discrepancy of 55%; B: Group B discrepancy of 56%; C: Group C discrepancy of 58%; D: Group D discrepancy of 58%.

In their study, O'Rahelly *et al*[18] observed a 40% reduction in the autopsy rate and other studies in the literature[9,19,20].

One factor that can have a positive influence on reducing the performance of hospital autopsies is the communication by the doctor of the importance of the autopsy to provide clarification of the cause of death[21], especially in sudden death in fetuses, infants and young people, from cardiac or non-cardiac causes and from genetic or non-cardiac causes.

Studies in the literature[22-25] report that male infants and children tend to have higher mortality rates than females, which may be explained by the greater vulnerability of males to perinatal complications, congenital diseases and genetic syndromes. In particular, male newborns are more likely to be born prematurely with low birth weight and to develop respiratory distress syndrome, as they have slower lung maturation than females, making them more susceptible to respiratory problems and consequently significantly increases the risk of neonatal mortality. Furthermore, another crucial aspect is the genetic vulnerability of males due to the presence of only one X chromosome compared to XX females, which makes them more susceptible by increasing mortality. In addition, it is important to underline that stillbirth represents a dramatic experience, not only for parents, but also for professionals, especially if it occurs in the last weeks of gestation.

Interestingly, in our case history, the major cause of death was pulmonary pathologies. This result may be due to the fact that lung pathologies are often subtle in their clinical manifestation and are among the pathologies that predominantly manifest themselves in long-term hospitalizations in patients hospitalized for another pathology, but also that the cause of death is often not clear and therefore clinicians are more sensitive to requesting a hospital autopsy. Therefore, it is clear that the hospital autopsy is still useful in the modern age to evaluate the clinical diagnostic accuracy. It is particularly important for fetuses and newborns to identify the various causes of death, both genetic and otherwise, and to be able to help future parents plan for subsequent pregnancies[26]. From a public health perspective, the autopsy can become a preventive tool for family members and the community and play a role in grieving[27].

Strategies to increase autopsy rates

Our study and others demonstrate that hospital autopsies have significantly decreased in recent years despite being a fundamental tool for understanding causes of death and for improving medical practices. Consequently, this reduction has resulted in a loss of valuable clinical information that could contribute to medical training, scientific research and improving the quality of care. Effective strategies to increase autopsy rates should involve raising awareness of healthcare personnel, patients and hospital policies.

A crucial first step is to invest in the continuous education and training of healthcare personnel, who must be constantly updated on the importance of autopsies and the procedures for requesting them through periodic refresher courses. The next step is empathetic communication by health professionals with families who are unaware of the benefits of autopsies, both for the medical community and for society. It is important to educate families and provide them with clear and easily understandable information through information leaflets and other resources that explain the autopsy process, its benefits and address ethical and religious issues.

Additionally, hospital policies are instrumental in providing guidelines and recommendations to establish mandatory autopsies in specific circumstances, such as in cases of sudden deaths.

CONCLUSION

In conclusion, the study of hospital autopsies at the Polyclinic of Bari shows that the discrepancy between clinical and autopsy diagnosis is 56.75%. Moreover, the hospital autopsy is still useful in the modern age, especially for the diagnosis of fetal and neonatal pathology who together account for 59% of autopsies, so that genetic and non-genetic diagnoses can be studied to help future parents for subsequent pregnancies. Focusing on the problems of stillbirth means ensuring adequate support for mothers and relatives, who are all too often left alone to face this traumatic event. Furthermore, analysis of hospital autopsy data over time reveals significant changes in patient demographics, medical specialties involved and causes of death. Despite these changes, the discrepancy between clinical and autopsy diagnoses remains an ongoing problem, underscoring the crucial importance of autopsies to improve diagnostic accuracy and the quality of medical care.

FOOTNOTES

Author contributions: Salzillo C and Basile R contributed equally; Salzillo C and Basile R contributed to the conceptualization and methodology; Salzillo C and Cazzato G performed the investigation; Basile R and Cazzato G were involved in data curation; Salzillo C, Basile R, and Cazzato G contributed to writing – original draft preparation; Marzullo A and Ingravallo G contributed to writing – review and editing; Marzullo A and Ingravallo G supervised the work; All authors have read and agreed to the published version of the manuscript.

Institutional review board statement: The data used in the document were obtained from autopsy reports in the database of the Pathology Unit, Department of Precision and Regenerative Medicine and Ionian Area, Bari Polyclinic, Italy. Access to the hospital archive for consultation of histopathological reports was approved and allowed by the director of the Pathology Unit. Furthermore, as this is a retrospective study of deceased patients, institutional review board approval is not required.

Informed consent statement: Data used in the paper were obtained from autoptic reports and do not need to be authorized for publication because they are considered anonymous and do not contain sensitive data. Moreover, according to Italian Law, the relatives of the deceased patients give their authorization to the use of the data for scientific purposes with the consent to the autopsy.

Conflict-of-interest statement: The authors declare no conflict of interest.

Data sharing statement: Not applicable.

Open-Access: This article is an open-access article that was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution NonCommercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <https://creativecommons.org/licenses/by-nc/4.0/>

Country of origin: Italy

ORCID number: Cecilia Salzillo 0009-0002-7531-3178; Gerardo Cazzato 0000-0003-0325-4316.

S-Editor: Liu JH

L-Editor: Filipodia

P-Editor: Zhao YQ

REFERENCES

- 1 **Cambiaghi M.** Andreas Vesalius (1514-1564). *J Neurol* 2017; **264**: 1828-1830 [PMID: 28303343 DOI: 10.1007/s00415-017-8459-2]
- 2 **Ghosh SK.** Giovanni Battista Morgagni (1682-1771): father of pathologic anatomy and pioneer of modern medicine. *Anat Sci Int* 2017; **92**: 305-312 [PMID: 27629485 DOI: 10.1007/s12565-016-0373-7]
- 3 **Prichard R.** Selected items from the history of pathology: karl von rokitansky (1804-1878). *Am J Pathol* 1979; **97**: 276
- 4 **Tan SY, Brown J.** Rudolph Virchow (1821-1902): "pope of pathology". *Singapore Med J* 2006; **47**: 567-568 [PMID: 16810425]
- 5 **Magnon R.** [Maurice Letulle (1853-1929)]. *Rev Infirm* 2009; **45** [PMID: 19317090]
- 6 **Sblano S, Arpaio A, Zotti F, Marzullo A, Bonsignore A, Dell'Erba A.** Discrepancies between clinical and autoptic diagnoses in Italy: evaluation of 879 consecutive cases at the "Policlinico of Bari" teaching hospital in the period 1990-2009. *Ann Ist Super Sanita* 2014; **50**: 44-48 [PMID: 24695252 DOI: 10.4415/ANN_14_01_07]
- 7 **Shojania KG, Burton EC.** The vanishing nonforensic autopsy. *N Engl J Med* 2008; **358**: 873-875 [PMID: 18305264 DOI: 10.1056/NEJMp0707996]
- 8 **Lundberg GD.** Low-tech autopsies in the era of high-tech medicine: continued value for quality assurance and patient safety. *JAMA* 1998; **280**: 1273-1274 [PMID: 9786381 DOI: 10.1001/jama.280.14.1273]
- 9 **Turnbull A, Osborn M, Nicholas N.** Hospital autopsy: Endangered or extinct? *J Clin Pathol* 2015; **68**: 601-604 [PMID: 26076965 DOI: 10.1136/jclinpath-2014-202700]
- 10 **McPhee SJ, Bottles K.** Autopsy: moribund art or vital science? *Am J Med* 1985; **78**: 107-113 [PMID: 3881019 DOI: 10.1016/0002-9343(85)90470-x]
- 11 **Scarl R, Parkinson B, Arole V, Hardy T, Allenby P.** The hospital autopsy: the importance in keeping autopsy an option. *Autops Case Rep* 2022; **12**: e2021333 [PMID: 35252044 DOI: 10.4322/acr.2021.333]
- 12 **Hamza A.** Perception of pathology residents about autopsies: results of a mini survey. *Autops Case Rep* 2018; **8**: e2018016 [PMID: 29780753 DOI: 10.4322/acr.2018.016]
- 13 **Esteban A, Fernández-Segoviano P.** The autopsy as a tool to monitor diagnostic error. *Intensive Care Med* 1999; **25**: 343-344 [PMID: 10342503 DOI: 10.1007/s001340050853]
- 14 **Xiao J, Krueger GR, Buja LM, Covinsky M.** The impact of declining clinical autopsy: need for revised healthcare policy. *Am J Med Sci* 2009; **337**: 41-46 [PMID: 19155753 DOI: 10.1097/MAJ.0b013e318184ce2b]
- 15 **Friberg N, Ljungberg O, Berglund E, Berglund D, Ljungberg R, Alafuzoff I, Englund E.** Cause of death and significant disease found at autopsy. *Virchows Arch* 2019; **475**: 781-788 [PMID: 31691009 DOI: 10.1007/s00428-019-02672-z]
- 16 **Veress B, Alafuzoff I.** A retrospective analysis of clinical diagnoses and autopsy findings in 3,042 cases during two different time periods. *Hum Pathol* 1994; **25**: 140-145 [PMID: 8119713 DOI: 10.1016/0046-8177(94)90269-0]
- 17 **Latten BGH, Kubat B, van den Brandt PA, Zur Hausen A, Schouten LJ.** Cause of death and the autopsy rate in an elderly population. *Virchows Arch* 2023; **483**: 865-872 [PMID: 37269366 DOI: 10.1007/s00428-023-03571-0]
- 18 **O'Rahelly M, McDermott M, Healy M.** Autopsy and pre-mortem diagnostic discrepancy review in an Irish tertiary PICU. *Eur J Pediatr* 2021; **180**: 3519-3524 [PMID: 34137920 DOI: 10.1007/s00431-021-04155-3]
- 19 **Lindström P, Janzon L, Sternby NH.** Declining autopsy rate in Sweden: a study of causes and consequences in Malmö, Sweden. *J Intern Med* 1997; **242**: 157-165 [PMID: 9279293 DOI: 10.1046/j.1365-2796.1997.00178.x]
- 20 **Latten BGH, Overbeek LIH, Kubat B, Zur Hausen A, Schouten LJ.** A quarter century of decline of autopsies in the Netherlands. *Eur J Epidemiol* 2019; **34**: 1171-1174 [PMID: 31728879 DOI: 10.1007/s10654-019-00572-9]
- 21 **Vignau A, Milikowski C.** The autopsy is not dead: ongoing relevance of the autopsy. *Autops Case Rep* 2023; **13**: e2023425 [PMID: 37292388 DOI: 10.4322/acr.2023.425]
- 22 **Ingemarsson I.** Gender aspects of preterm birth. *BJOG* 2003; **110** Suppl 20: 34-38 [PMID: 12763109 DOI: 10.1016/s1470-0328(03)00022-3]
- 23 **Rysavy MA, Horbar JD, Bell EF, Li L, Greenberg LT, Tyson JE, Patel RM, Carlo WA, Younge NE, Green CE, Edwards EM, Hintz SR, Walsh MC, Buzas JS, Das A, Higgins RD; Eunice Kennedy Shriver National Institute of Child Health and Human Development Neonatal Research Network and Vermont Oxford Network.** Assessment of an Updated Neonatal Research Network Extremely Preterm Birth Outcome Model in the Vermont Oxford Network. *JAMA Pediatr* 2020; **174**: e196294 [PMID: 32119065 DOI: 10.1001/jamapediatrics.2019.6294]
- 24 **Garite TJ, Combs CA, Maurel K, Das A, Huls K, Porreco R, Reinsner D, Lu G, Bush M, Morris B, Bleich A; Obstetrix Collaborative Research Network.** A multicenter prospective study of neonatal outcomes at less than 32 weeks associated with indications for maternal admission and delivery. *Am J Obstet Gynecol* 2017; **217**: 72.e1-72.e9 [PMID: 28267444 DOI: 10.1016/j.ajog.2017.02.043]
- 25 **Tietzmann MR, Teichmann PDV, Vilanova CS, Goldani MZ, Silva CHD.** Risk Factors for Neonatal Mortality in Preterm Newborns in The Extreme South of Brazil. *Sci Rep* 2020; **10**: 7252 [PMID: 32350375 DOI: 10.1038/s41598-020-64357-x]
- 26 **Xu Y, Cheng C, Zheng F, Saiyin H, Zhang P, Zeng W, Liu X, Liu G.** An audit of autopsy-confirmed diagnostic errors in perinatal deaths: What are the most common major missed diagnoses. *Heliyon* 2023; **9**: e19984 [PMID: 37809936 DOI: 10.1016/j.heliyon.2023.e19984]
- 27 **McPhee SJ, Bottles K, Lo B, Saika G, Crommie D.** To redeem them from death. Reactions of family members to autopsy. *Am J Med* 1986; **80**: 665-671 [PMID: 3963043 DOI: 10.1016/0002-9343(86)90822-3]



Published by **Baishideng Publishing Group Inc**
7041 Koll Center Parkway, Suite 160, Pleasanton, CA 94566, USA

Telephone: +1-925-3991568

E-mail: office@baishideng.com

Help Desk: <https://www.f6publishing.com/helpdesk>

<https://www.wjgnet.com>

