

## Study The Relation Between Fingerprint Patterns And Cancer (Breast & Prostate)

Ritu Sharma<sup>1</sup>, Dr. Ashok Sagar<sup>2\*</sup>, Ms. Neha Jha<sup>3</sup>, Dr. Mukesh Kumar<sup>4</sup>, Dr. Amit Singhal<sup>5</sup>, Dr Shri Bhagwan<sup>6</sup> Jay prakash singh Rajput<sup>7</sup>

<sup>1</sup>M.sc Student Forensic Science, Department of Forensic Science, Santosh Deemed to be University, Ghaziabad, India, ritukaushik2201@gmail.com

<sup>2</sup>Professor and Head of Department, Department of Forensic Medicine & Toxicology, Santosh Deemed to be University, Ghaziabad, India. Email: dr.sagarashok@gmail.com

<sup>3</sup>Senior Tutor, Department of Forensic Science, Santosh Deemed to be University, Ghaziabad, India, Nehajha524@gmail.com

<sup>4</sup>Associate Professor Department, Department of Forensic Medicine & Toxicology, Santosh Deemed to be University, Ghaziabad, India. Email: mukeshudct1877@gmail.com

<sup>5</sup>Professor, Department of Pharmacology, Maharishi Markndeswar Institute of Medical Sciences and Research Mullana, Ambala, Haryana, India. Email: dramitkmmc@gmail.com

<sup>6</sup>Professor, Amrita school of medicine, Faridabad, shribhagwan315@gmail.com

<sup>7</sup>Associate Professor, Department of Physiology, Netaji Subash Medical College and Hospital, Bihta, Patna, jpsrajput50@gmail.com

---

### ABSTRACT

**Introduction:** Dermatoglyphics means "skin patterns" and is a way of studying the tiny lines and shapes on the skin of fingers, palms, toes, and soles. Fingerprint pattern can serve as indicators for genetically linked health conditions. People with certain health conditions or syndromes often have different patterns on their fingertips compared to others. These patterns form during the fifth and sixth weeks of a baby's life and are fully developed by the 21st week. Once they are formed, they never change. These lines help fingers grip things and are also used to identify people in a unique way.

**Aim:** To compare fingerprint patterns among breast cancer patients, prostate cancer patients and healthy controls to identify potential fingerprint markers associated with cancer predisposition.

**Methods:** This was a comparative and case-control study set up for a comparison between two groups to observed the patterns on people's fingers, called dermatoglyphics, in those with breast or prostate cancer compared to people who are healthy. Total of 70 people took part in this research study. 35 people who had been diagnosed with breast/prostate cancer. Additionally, 35 healthy individuals, matched for age and sex with cancer patients and had no history of cancer were took for study. Standard ink-rolled ten-digit fingerprints were recorded.

**Results:** Among (35) healthy controls, Loops were identified as the predominant fingerprint pattern (64.2%), with Arches (20.6%) and Whorls (15.2%). However, in (30) breast cancer patients there was a marked shift in pattern distribution. In this group, whorls emerged as significantly more common, present (45%) establishing them as the most frequent pattern. Loops showed a slight decrease, found (40%), while Arches were observed (15%). This inversion in the dominant pattern suggests a possible link between the increased presence of Whorls and the occurrence of breast cancer. In a similar vein, among (5) prostate cancer patients, Whorls were more identified as the most prevalent pattern, observed (48%) and Loops were present (36%) and Arches were the least common about (16%).

**Conclusion:** These findings suggest that the heightened occurrence of whorl patterns and a diminished count of loop patterns, notably observed in cases of breast cancer, may function as potential indicators for screening or assessing individual risk. Although the fingerprints alone are not a unique diagnostic indicator; they do represent a non-invasive and low-cost method of investigation.

**Keywords:** Fingerprints; breast cancer; prostate cancer; whorl pattern; cancer biomarkers.

**How to cite this article:** Sharma R, Sagar A, Jha N, Kumar M, Singhal A, Bhagwan S, Rajput JPS. Study The Relation Between Fingerprint Patterns And Cancer (Breast & Prostate). Int J Drug Deliv Technol. 2026;16(58s): 433-438. DOI: 10.25258/ijddt.16.58s.42

**Source of support:** Nil.

**Conflict of interest:** None.

---

### INTRODUCTION

Dermatoglyphics means "skin patterns" and is a way of studying the tiny lines and shapes on the skin of fingers, palms, toes, and soles. Fingerprint pattern can serve as indicators for genetically linked health conditions. People with certain health conditions or syndromes often have different patterns on their fingertips

\*Author for Correspondence: dr.sagarashok@gmail.com

compared to others. These patterns form during the fifth and sixth weeks of a baby's life and are fully developed by the 21st week. Once they are formed, they never change. These lines help fingers grip things and are also used to identify people in a unique way. Doctors can use these patterns for medical tests without taking any samples. In 1892, Francis Galton & Edward Henry

began their 1<sup>st</sup> study on fingerprints & classified them. They described three main types of fingerprint patterns: Arches, Loops and Whorls [7,8].The patterns include fingerprints, ridge patterns & palm prints. By the passage of time, studying fingerprints has given people a lot of knowledge about this study. Palm prints have more information than fingerprints.

In 1892, Francis Galton & Edward Henry conducted pioneering research on fingerprints, establishing their classification system. They categorized the fingerprints into 3 main types: Arches, Loops and Whorls. Archaeologists found fingerprints on pottery tablets in Babylon that are more than 4,000 years old, tracking back to 1792–1750 BCE.

- (1) Arches pattern has two kinds: (a) Plain Arch & (b) Tented Arch.
- (2) The Loop pattern divided into two types: (a) Radial Loops & (b) Ulnar Loops.
- (3) The Whorls pattern have four subgroups:(a) Plain Whorl, (b)Central Pocket Loop, (c) Double Loop, and (d) Accidental Whorl.

#### **Link between cancer (breast & prostate) and fingerprints**

Cancer is a major global contributor to mortality, accounting for a significant number of deaths worldwide. Some studies suggest that looking at fingerprint patterns might help spot people who could be more likely to get certain genetic diseases early [14]. Because of this, this study looks at existing research on different types of hand fingerprint patterns and whether they might be connected to a higher risk of cancer. The study aims to discover which fingerprint patterns are correlated with cancer and the corresponding cancer types.

#### **MATERIAL & METHODS**

##### **Study Population**

This was a **comparative and case-control study** set up for a comparison between two groups to observed the patterns on people's fingers, called dermatoglyphics, in those with breast or prostate cancer compared to people

who are healthy. Total of 70 people took part in this research study-35 people who had been diagnosed with breast/prostate cancer. Cancer patients were recruited from Santosh Hospital Ghaziabad and Subharthi Hospital in Meerut.

Additionally, 35 healthy individuals, matched for age and sex with cancer patients and had no history of cancer were took for study. Those healthy people selected from Forensic Medicine & Toxicology Department (Santosh deemed to be University).

##### **Inclusion Criteria**

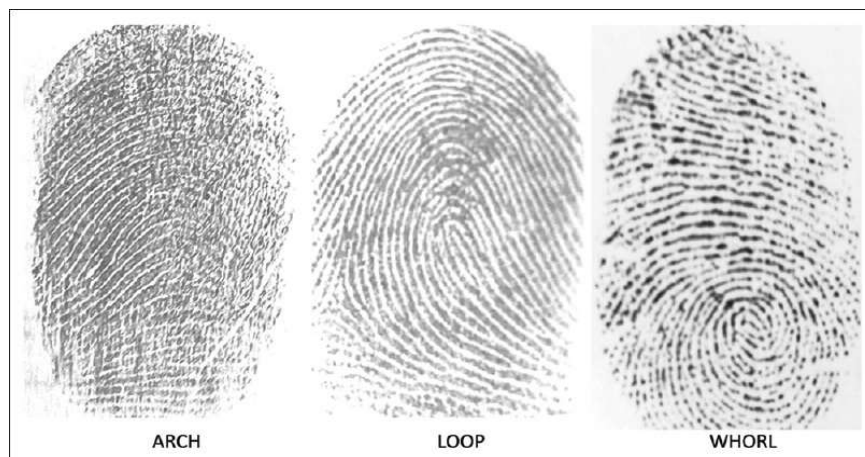
Diagnosed cases of breast or prostate cancer.

##### **Exclusion Criteria**

Finger deformities or dermatological conditions. Hand injuries or scars affecting ridge clarity.

##### **Data collection & analysis**

Hand hygiene was ensured by asking each patient to cleanse their hands using any soap and water before collection of data. This helps make the fingerprints clear. They then dried their hands with a clean towel or paper. Next, a thin layer of ink was spread over a flat surface using the metal roller. The roller helped make sure the ink was spread evenly without making it too thick, which could cause smudges. Then, each finger was rolled gently from the tip of the nail to the other end across the inked surface and then onto the fingerprint card. The rolling was done from the thumb side (radial) to the little finger side (ulnar) to cover all the lines properly. This was repeated for all ten fingers, starting with the right hand, from the thumb to the little finger, and then the left hand in the same order. Each card was labelled with the participant's name, contact number and signed consent form and the date when the fingerprints were taken, and whether it was the left or right hand. After collecting the fingerprints, the cards were left to dry in the air for 10 to 15 minutes to prevent them from getting smudged. Lastly, all fingerprints were observed under 10x magnifying lens to analysed the ridge details, core, delta and different types of fingerprint patterns (Arches, loops and Whorls).



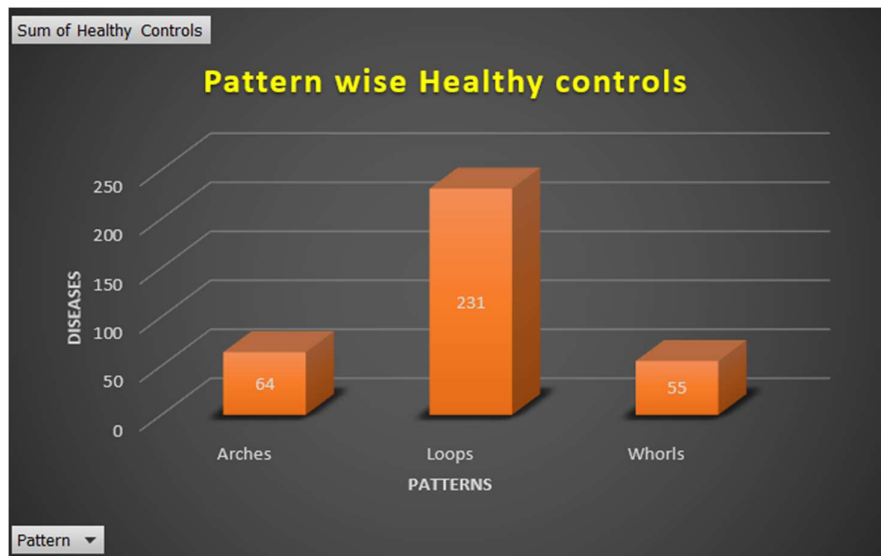
#### **RESULT**

The study aimed to investigate the distribution of dermatoglyphic patterns—Loops, Whorls, and

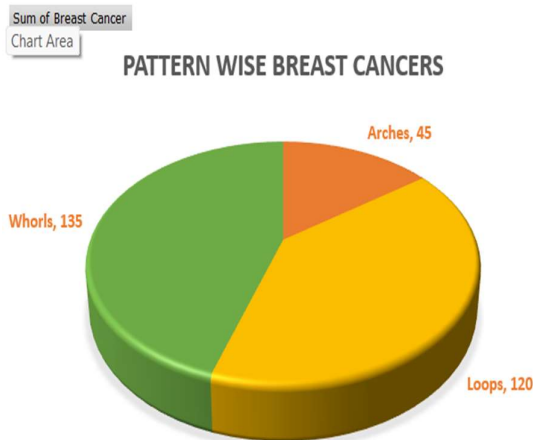
Study The Relation Between Fingerprint Patterns And Cancer (Breast & Prostate)

Arches—among healthy individuals, breast cancer patients, and prostate cancer patients. Among 350 patterns of **healthy controls (35)**, **Loops were identified as the predominant fingerprint pattern**, accounting for 231 cases (64.2%), with Arches appearing in 74 instances (20.6%) and Whorls in 55 instances (15.2%). This observed pattern aligns with the general population's typical distribution, characterized by the prevalence of loops. However, in **(30) breast cancer patients (n = 300)**, there was a marked shift in pattern distribution. In this group, **whorls emerged as significantly more common**, present in 135 instances (45%) establishing them as the most frequent pattern. Loops showed a slight decrease, found in 120 instances (40%), while Arches were observed in only 45 cases (15%). This inversion in the dominant pattern suggests a possible link between the increased presence of

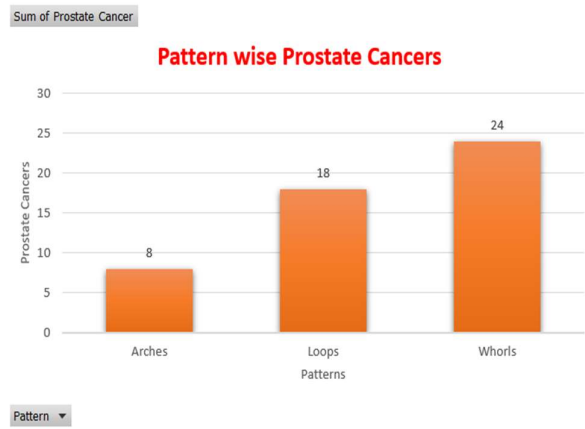
Whorls and the occurrence of breast cancer. In a similar vein, among 5 prostate cancer patients (n=50), **Whorls were more identified as the most prevalent pattern**, observed in 24 fingers (48%). Loops were present in 18 fingers (36%), and Arches were the least common, found in only 8 fingers (16%). This pattern mirrors the trend observed in breast cancer patients, further supporting the hypothesis that Whorls may be more prevalent in individuals with certain types of cancer. Even though Arches have always been the least common pattern across all three groups, their reduction in cancer patients is a component of the particular pattern shift. Fingerprint pattern analysis shows promise as a supportive, non-invasive screening method for early cancer detection or risk evaluation; however, further investigation using broader and more diverse population studies is necessary.



(Figure.1)



(Figure.2)



(Figure.3)

• In healthy control subjects, loop patterns were most prevalent (66%), followed by Arches (18.3%) and then Whorls (15.7%).(Figure.1)

• Breast Cancer Patients had a higher prevalence of whorls (45%), followed by loops (40%), and arches (15%).(Figure.2)

## Study The Relation Between Fingerprint Patterns And Cancer (Breast & Prostate)

• Whorl patterns were more common in prostate cancer patients (48%) than loop (36%) or Arch (16%) patterns.(Figure.3)

The patterns were compared between healthy people and those diagnosed with either breast or prostate

Group	Loops (L)	Whorls (W)	Arches (A)	Total Prints
Healthy Controls	231	55	64	350
Breast Cancer Patients	120	135	45	300
Prostate Cancer Patients	18	24	8	50

### DISCUSSION

The current study showed a distinct difference in the distribution of fingerprint patterns between healthy subjects and patients with breast or prostate cancer. Loop fingerprint pattern was most likely observed in healthy group of control sample nearly (66%), and this finding is consistent with the dermatoglyphic literature. In patients diagnosed with breast cancer, the most common type of fingerprint was whorl (45% frequency). Whorls were also the most frequently identified fingerprint type in prostate cancer patients (48%), which is a significant difference compared to a healthy group or controls. The altered frequency of patterns suggests an association between the morphology of fingerprints and some predisposition to and pattern of disease. The predominant presence of whorl patterns in cancer patients, contrasted with their relative rarity in healthy individual's fingerprints (15.7%), highlights a significant difference that warrants further investigations and important questions related to genetic factors and developmental aesthetic can be fundamentally assessed as indicators of cancer or potential cancer conditions.

### CONCLUSION

This research examined the distribution of various fingerprint patterns: loops, whorls, and arches in healthy individuals, and patients with breast or prostate cancers. The findings are markedly different in dermatoglyphics patterns in the control group when compared to the cancer samples. While loops were the most common fingerprint pattern in healthy individuals (66%), their prevalence decreased in patients with breast cancer (40%) and prostate cancer (36%).

A considerably higher number of whorl fingerprint pattern were mostly observed in female patients having breast cancer (45%) in comparison with control sample (16%), suggesting a possible link to the disease. Arches were somewhat stable across the groups, but somewhat more prevalent in the healthy individuals.

These findings could mean that the fingerprint patterns reflect biological changes, minor or otherwise, that are somehow caused or associated to cancer by genetic and or developmental means.

The heightened occurrence of whorl patterns and a diminished count of loop patterns, notably observed in cases of breast cancer, may function as potential indicators for screening or assessing individual risk. Although the fingerprints alone are not a unique

cancer. The partitioning of loops, whorls, and arches can be accounted for in each grouping as demonstrated in the following table:

diagnostic indicator; they do represent a non-invasive and low-cost method of investigation.

There is great potential with dermal ridges going forward; if these patterns are repeated in further studies, we can begin to understand and predict actions. Further research involving larger and more varied group is necessary to analyse the internal patterns and determine their potential cause.

Consequently, dermatoglyphics offers a potentially valuable yet underutilized avenue for enhancing personalized cancer patient risk assessment.

### DECLARATION

#### Funding-

No external funding was received.

#### Conflicts of Interest-

The authors declare no conflict of interest.

#### Ethical Approval-

Approved by Institutional Human Ethics Committee (F. No. SU/2025/CRF/218).

#### Informed Consent-

Written informed consent was obtained from all participants.

### REFERENCES

- Seltzer MH, Plato CC, Engler PE, Fletcher HS. Digital dermatoglyphics and breast cancer. *Breast Cancer Res Treat.* 1982; 2:261–65
- Chintamani, Khandelwal R, Mittal A, Saijanani S, Tuteja A, Bansal A, Bhatnagar D, Saxena S. Qualitative and quantitative dermatoglyphic traits in patients with breast cancer: a prospective clinical study. *BMC Cancer.* 2007 Mar 13; 7:44.
- Raizada A, Johri V, Ramnath T, Chowdhary DS, Garg RP. A cross-sectional study on the palmar dermatoglyphics in relation to carcinoma breast patients. *J Clin Diagn Res.* 2013 Apr;7(4):609–12.
- Sridevi NS, Silvia CRW, Kulkarni R, Seshagiri C. Palmar dermatoglyphics in carcinoma breast of Indian women. *Rom J Morphol Embryol.* 2010;51(3):547–50.
- Sukre SB, Laeeque M, Mahajan A, Shewale SN. Dermatoglyphics in the identification of women either with or at risk for breast cancer. *Int J Basic Med Sci.* 2012;3(5):160–65.
- Musanović J, Metović A, Pepić E, et al. Predictive values of quantitative analysis of finger and palmar dermatoglyphics in patients with breast cancer for Bosnian-Herzegovinian population. *J Evol Med Dent Sci.* 2018;7(24):2855–60.

7. Oladipo GS, Paul CW, Bob-Manuel IF, Fawehinmi HB, Edibamode EI. Study of digital and palmar dermatoglyphic patterns of Nigerian women with malignant mammary neoplasm. *J Appl Biosci*. 2009; 15:829–34.
8. Gupta A, Karjodkar FR. Role of dermatoglyphics as an indicator of precancerous and cancerous lesions of the oral cavity. *Contemp Clin Dent*. 2013 Oct;4(4):448–53.
9. Dodia VS, Odedra SP, Shah KH, Monpara PC, Vyas PM, Pillai JP. The association of fingerprint patterns with oral potentially malignant disorders and oral cancer: a dermatoglyphic study. *J Oral Maxillofac Pathol*. 2022 Jul-Sep;26(3):420.
10. Tonkaboni A, Etemadian M, Manifar S, Shirkhoda M, Gharehdaghi J, Kharazi Fard MJ. Investigating the relationship between fingerprint pattern and development of oral squamous cell carcinoma. *J Dent (Shiraz)*. 2022 Jun;23(2):144–50.
11. Venkatesh E, et al. Palmar dermatoglyphics in oral leukoplakia and oral squamous cell carcinoma patients. *J Indian Acad Oral Med Radiol*. 2008;20(3):94.
12. Jatti D, Kantraj YDB, Nagaraju R. Role of dermatoglyphics in malignant and potentially malignant disorders of the oral cavity: a cross-sectional study. *J Indian Acad Oral Med Radiol*. 2014;26(4):379.
13. Oladipo GS, et al. Dermatoglyphics of prostate cancer patients. *Curr Res J Biol Sci*. 2009;1(3):131–34.
14. Ganorkar DY, Kawale DS, Pandit DSV, Kumar Jha DR. Study of dermatoglyphics in thalassemia major... (not cancer but included for pattern methodology). *IJSR*. 2016;15(08):79–84.
15. Paul G, Kalyan GS, Kaur H, Chhabra U. Dermatoglyphic pattern on fingertips of thalassemia patients: a case control study. *IJAR*. 2018;6(2):5322–38.
16. Mahato LO, Naidu SS, Rangari KA, Rahule AS, Bashir MS. Comparative study of fingerprint pattern in thalassemic population. *Medico-Legal Update*. 2013;13(1):40–44.
17. Wijerathne BT, Meier RJ, Agampodi SB. Dermatoglyphics in hypertension and kidney diseases: implications for genetic predisposition. *J Physiological Anthropology*. 2015
18. Fuller IC. Inherited Predisposition to Cancer? A dermatoglyphic study. *Br J Cancer*. 1973; 28:186–89.
19. “The role of hand fingerprints on predisposition of cancer development” — review of associations between fingerprint patterns and various cancers including gynecological, oral, gastric, prostate, leukemia, pituitary tumors. [Elsevier review] 2023.
20. Kahn HS, Ravindranath R, Valdez R, et al. Early prenatal fingerprint markers and disease predisposition: insights from developmental epidemiology. *Am J Hum Biol*. 2008;20(1):59–65. (gestational fingerprint markers for later disease risk).
21. He J, Martinez-Lozano Sinues P, Hollmén M, Li X, Detmar M, Zenobi R. Fingerprinting breast cancer vs normal mammary cells by mass spectrometric analysis of volatiles. *Sci Rep*. 2014;4:5196.
22. Ferguson D, Kroeger-Lui N, Dreisbach D, Hart CA, Sanchez DF, Oliveira P, et al. Full fingerprint hyperspectral imaging of prostate cancer tissue microarrays within clinical timeframes using quantum cascade laser microscopy. *Analyst*. 2025; 150:1741–1753.
23. Fairey A, et al. Clinical analysis of EV-Fingerprint to predict grade group 3 and above prostate cancer and avoid prostate biopsy. *Cancer Med*. 2023.
24. Snider AJ, Seeds MC, Johnstone L, Snider JM, Hallmark B, Dutta R, et al. Identification of plasma glycosphingolipids as potential biomarkers for prostate cancer (PCa) status. *Biomolecules*. 2020;10(10):1393.
25. Anonymous (UCL Cancer Institute). Prostate cancer ‘fingerprint’ detected in blood sample. *J Clin Invest*. 2020 Mar 10.
26. He Ahr A, Karn T, Solbach C, Seiter T, Strebhardt K, Holtrich U, et al. Identification of high-risk breast-cancer patients by gene expression profiling. *Lancet*. 2002;359(9301):131–132.
27. van de Vijver MJ, He YD, van’t Veer LJ, Dai H, Hart AA, Voskuil DW, et al. A gene-expression signature as a predictor of survival in breast cancer. *N Engl J Med*. 2002;347(25):1999–2009.
28. Sotiriou C, Wirapati P, Loi S, Harris A, Fox S, Smeds J, et al. Gene expression profiling in breast cancer: understanding the molecular basis of histologic grade to improve prognosis. *J Natl Cancer Inst*. 2006;98(4):262–272.
29. Kunthoth S, Al-Maadeed S, Akbari Y, Al Saady R. Computational methods for breast cancer molecular profiling through routine histopathology: a review. *arXiv*. 2024 Dec;2412.10392.
30. Nassif AB, Talib MA, Nasir AM, Afadar Y, Elgendy O. Breast cancer detection using artificial intelligence techniques: a systematic literature review. *arXiv*. 2022 Mar;2203.04308.
31. Zhou J, Rhrissorrakrai K. Discovering multi-omic biomarkers for prostate cancer severity using machine learning. *arXiv*. 2024 Oct;2410.22387.
32. Fard AT, Srihari S, Ragan MA. An evaluation of DNA-damage response and cell-cycle pathways for breast cancer classification. *arXiv*. 2015 May;1505.04846.
33. Davis et al. Molecular biology underlying metastatic PCa and Decipher genomic test for fingerprinting. In: *Biology of the Cell*. 2015.
34. (Additional) Tipatet KS, Hanna K, Davison-Gates L, Kerst M, Downes A. Subtype-Specific Detection in Stage Ia Breast Cancer: integrating Raman spectroscopy, machine learning, and liquid biopsy for personalised diagnostics. *J Biophotonics*. 2025;18: e202400427.
35. (Supplement) AI-powered blood test spots earliest breast cancer signs combining Raman spectroscopy with machine learning. Pilot stage accuracy ~98 % at stage Ia. *J Biophotonics*. 2025.

36. (Supplement) AI detects prostate cancer tumour margins more accurately than doctors by integrating imaging/biopsy/biomarker data. *Eur Urol*. 2023 publication.
37. Rotten supplementary on radiogenomics mapping molecular imaging fingerprint in prostate and breast cancer visibility. *Radiology, Eur Urol Open Science*. 2018–2020.
38. Farha MW, Salami SS. Biomarkers for prostate cancer detection and risk stratification. *Ther Adv Urol*. 2022; 14:17562872221103988.