

Cross-Sensor Micro-Texture Material Classification and Smartphone Acquisition do not go well together

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Abstract

Intrinsic, non-invasive product authentication is still an important topic as it does not generate additional costs during the production process. This topic is of specific interest for medical products as non-genuine products can directly effect the patients' health. This work investigates micro-texture classification as a mean of proving the authenticity of zircon oxide blocks (for dental implants). Samples of three different manufacturers were acquired using four smartphone devices with a clip-on macro lens. In addition, an existing drug packaging material database was utilized. While the intra-sensor micro-texture classification worked well, the cross-sensor classification results were less promising. In an attempt to track down the limiting factors, intrinsic sensor features usually used in device identification were investigated as well.

Main Results

- Intra-Sensor Material Classification works well
- Cross-Sensor Material Classification does not work so well
- Sensor Classification works better than Material Classification
- Location dependent parts of PRNU are not the reason why sensor classification performance is better

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Introduction

Motivation

- Counterfeit products are a significant problem: Causing economic damage to the manufacturers
- Especially in medical/health care applications have direct influence on patients' health
- Medical and health related products usually high priced - hence of higher interest for product forgers
- Different ways to establish the authenticity of products
 - Intrinsic: based on the product's properties, no external markers etc. needed
 - Extrinsic: based on QR codes, external markers, validated supply chains, etc.

Goals of this work:

- Evaluate effect of scaling in cross sensor scenario for existing drug package database
- Scope of existing database:
 - 3 capturing devices
 - 5-6 drug manufacturers
 - 3 modalities (cardboard, blister top, blister bottom)
- Acquisition of zircon oxide block ceramic database
 - zircon oxide blocks from 3 manufacturers
 - 4 smartphone devices
- Can we classify ceramic manufacturers based on samples? (Intra-Sensor & Cross-Sensor)

Texture Classification Pipeline

- Different feature extraction schemes:
 - Dense SIFT (SIFT)
 - Dense Micro-block Difference (DMD)
 - Local Binary Pattern (LBP)
 - Local Phase Quantization (LPQ)
 - Weber Pattern (WP)
- Followed by a PCA based dimensionality reduction, a Fisher vector encoding and finally an SVM based classification

Ceramic Data Acquisition Setup

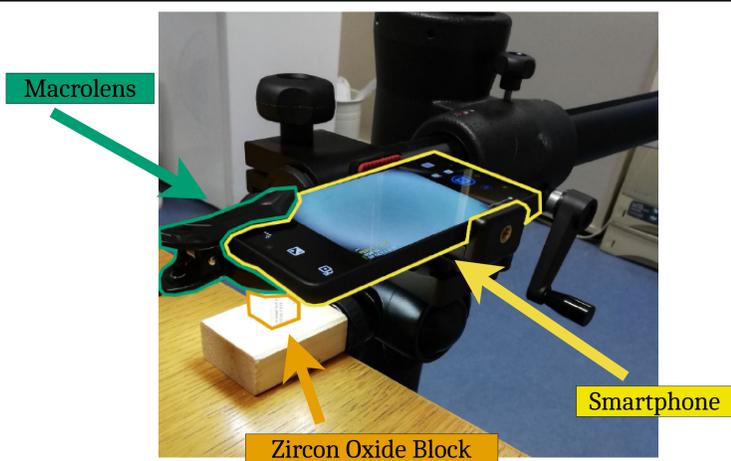


Figure 1: Image acquisition setup for zircon oxide blocks.

Ceramic Data Patching

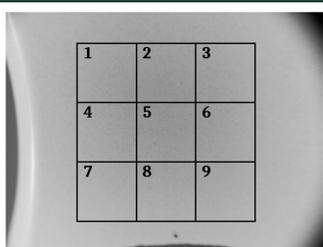


Figure 2: Acquired image with distortions & patching strategy.

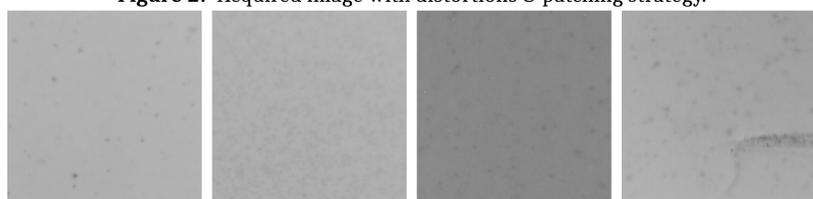


Figure 3: Ceramic data sample patches.

Samples from (pre-existing) Drug Package Database

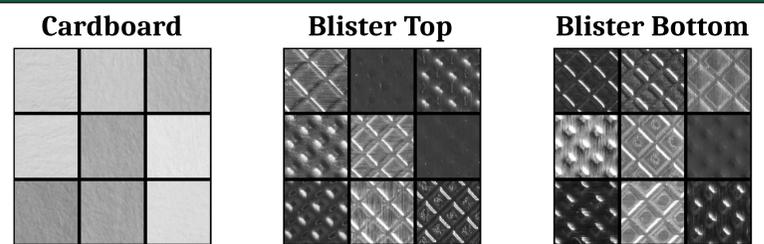


Figure 4: Drug data sample patches for different modalities.

Experiment 1: Intra-Sensor Material Classification

	SIFT	DMD	LBP	WP	LPQ
Cardboard	90.18	90.14	70.21	61.09	31.78
Blister Top	99.89	99.30	89.98	74.17	38.48
Blister Bottom	99.36	97.88	69.92	56.12	24.26
Ceramic	98.88	97.46	72.90	71.04	51.60

Table 1: Intra-sensor material classification. Averaged accuracy.

Experiment 2a: Cross-Sensor Ceramic Data Material Classification

	SIFT	DMD	LBP	WP	LPQ
Ceramic	68.60	60.74	36.17	55.65	31.13

Table 2: Leave-one-sensor-out ceramic material classification. Averaged accuracy.

Experiment 2b: Cross-Sensor Drug Package Data Scaling

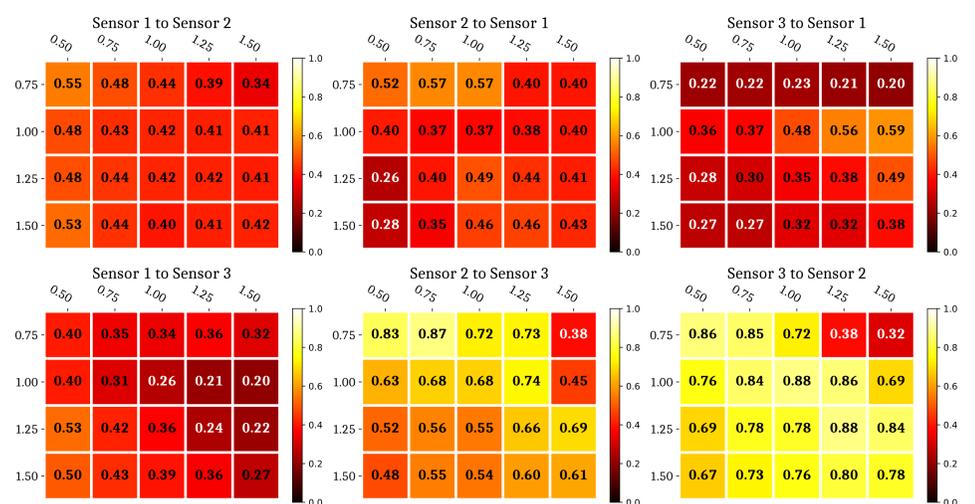


Figure 5: Different scale factors on the drug data (blister top-side) using SIFT.

Experiment 3: Sensor Classification

	SIFT	DMD	LBP	WP	LPQ
Cardboard	100.0	100.0	99.75	99.41	75.48
Blister Top	99.42	99.29	81.01	92.95	81.86
Blister Bottom	98.66	98.63	69.66	86.15	73.08
Ceramic	100.0	99.92	98.66	92.44	83.55

Table 3: Switch classes (manufacturers) and devices = Sensor identification. Averaged accuracy.

Experiment 4: Influence of PRNU

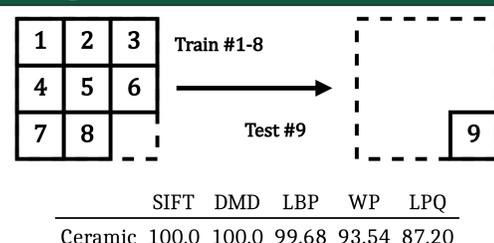


Table 4: Evaluation influence of PRNU, through testing a patch from location excluded in test data. Averaged accuracy.