

25 **Abstract**

26 A novel nanocomposite-inspired *in-situ* broadband ultrasonic sensor previously developed,
27 with carbon black as the nanofiller and polyvinylidene fluoride as the matrix, was
28 networked for acousto-ultrasonic wave-based passive and active structural health
29 monitoring (SHM). Being lightweight and small, this kind of sensor was proven to be
30 capable of perceiving strain perturbation in virtue of the tunneling effect in the formed
31 nanofiller conductive network when acousto-ultrasonic waves traverse the sensor.
32 Proof-of-concept validation was implemented, to examine the sensor performance in
33 responding to acousto-ultrasonic waves in a broad frequency regime: from acoustic
34 emission (AE) of lower frequencies to guided ultrasonic waves (GUWs) of higher
35 frequencies. Results have demonstrated the high fidelity, ultrafast response and high
36 sensitivity of the sensor to acousto-ultrasonic waves up to 400 kHz yet with an ultra-low
37 magnitude (of the order of micro-strain). The sensor is proven to possess sensitivity and
38 accuracy comparable with commercial piezoelectric ultrasonic transducers, whereas with
39 greater flexibility in accommodating curved structural surfaces. Application paradigms of
40 using the sensor for damage evaluation have spotlighted the capability of the sensor in
41 compromising “sensing cost” with “sensing effectiveness” for passive AE- or active
42 GUW-based SHM.

43

44 **Keywords:** ultrasonic sensor; acousto-ultrasonics; nanocomposite sensor; guided ultrasonic
45 waves; acoustic emission; structural health monitoring

46

47 **1. Introduction**

48 Acousto-ultrasonics, a coalescence of acoustic emission (AE) and ultrasonic
49 characterization [1, 2], has been a subject of intense scrutiny over the years and is now on the
50 verge of maturity for real-world engineering applications. It has gained prominence for
51 developing diverse structural health monitoring (SHM) approaches [3-11] in the past decade,
52 and central to the increased preference of using acousto-ultrasonics lies preliminarily in the
53 fact that it exploits the merits from both AE and guided ultrasonic waves (GUWs) in a broad
54 frequency regime, enabling monitoring at multi-scale so as to accommodate different
55 demands. Acousto-ultrasonics is demonstrably cost-effective in striking a compromise
56 among resolution, detectability, practicality and cost, corroborating the philosophy of *in-situ*
57 SHM.

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59 As a highly sophisticated technique, acousto-ultrasonics-based SHM relies heavily on
60 integrated sensor networks, real-time digital signal processing and pattern recognition – a
61 feature that distinguishes it from conventional non-destructive evaluation. In particular,
62 “sensing”, via a sensor network, plays the most rudimentary yet critical role, to collect
63 ambient information and system parameters, whereby a perception on the health status of an
64 inspected structure can be developed. The sensors to serve acousto-ultrasonics-based SHM
65 have been explored extensively [8, 12-21], as typified by piezoelectric wafer active sensors
66 (PWAS) [8, 22], SMART Layer[®] [17], polyvinylidene fluoride (PVDF)-based interdigital
67 transducers (IDTs) [16], macro fiber composites (MFCs) [18], and circular sensing ring [19]
68 to name a few. In common practice, a limited number of sensors are spatially distributed and
69 networked, to meet a minimal threshold guaranteeing the adequacy of information. During
70 network configuration, a paramount target is to balance “sensing cost” with “sensing
71 effectiveness” – to acquire adequate information with the least weight/volume penalty to the

72 inspected structure. However, owing to its distributed nature, a distributed sensor network
73 may “overlook” certain information, leading to inaccurate or even erroneous results. This
74 has entailed continued efforts to attempt new breeds of sensors for acousto-ultrasonics-based
75 SHM, in lieu of the conventional types of sensor.

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77 Remarkable advances and technological breakthroughs in nanotechnology, materials science,
78 manufacturing and electronic packaging have ushered in a new avenue to develop novel
79 sensors that are inspired by nanocomposites [23-39]. Using a variety of nanofillers such as
80 carbon black (CB) [28, 40], carbon fiber (CF) [41, 42], carbon nanotube (CNT) [31-33] or
81 graphene [43, 44], a nanocomposite sensor can be endowed with merits such as low density,
82 excellent flexibility, easy tailorability in shape and size, chemical stability, low fabrication
83 cost, along with good electrical and mechanical performance. The use of nanocomposite
84 sensors can now be found in a wide application domain, exemplarily including damage
85 identification [23-26], strain measure [27-37, 43, 44] and gas leak detection [38-40, 45, 46].

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87 Nevertheless, the development and use of nanocomposite sensors for
88 acousto-ultrasonics-based SHM is fairly lacking, because of the bottleneck that most
89 nanocomposite sensors may have: the measurand – the change in electrical conductivity of
90 the nanofiller-formed conductive network – is a global indicator with a uniform value
91 between a pair of electrodes, and it reflects only the holistic alteration in material
92 properties right underneath the sensor patch. Limited by this, this type of sensor is unable
93 to respond to damage away from the sensor, and distinguish different locations and degrees
94 of severity of damage. In addition, in order to perceive damage-induced change reliably,
95 especially the ultralow strain magnitude (of the order of micro-strain) in an ultrasonic

96 frequency regime, the sensitivity (gauge factor) of the nanocomposite sensors is to be
97 improved.

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99 In an earlier study [28], the authors have developed and fabricated a novel nanocomposite
100 sensor (made of CB and PVDF). Proven the first of nanocomposite sensor capable of
101 perceiving ultrasonic signals, the developed sensor sees the potential of being applied to
102 acousto-ultrasonic-based SHM. Compared with single-phase rigid ceramic sensor (*e.g.*, lead
103 zirconate titanate (PZT)), the developed nanocomposite sensors, being flexible and
104 lightweight, shows more adaptability to curved structures with a possibility of dense sensor
105 deployment. And compared with metallic foil-based strain gauges, the developed
106 nanocomposite sensor offers a higher gauge factor [28] and therefore greater sensitivity to
107 structural or ambient changes and higher possibility of sensing faint elastic disturbance
108 induced by acousto-ultrasonics. Nevertheless, the quantitative evaluation of the sensing
109 performance of the developed sensor for its applications to SHM has yet been fully
110 demonstrated. Relevant issues associated with the application, including parameter setup,
111 sensor networking, measurement system configuration, algorithm development, etc., are to
112 be explored.

113

114 In recognition of the stated deficiencies of nanocomposite sensors when they are attempted
115 to accommodate acousto-ultrasonics-based SHM, this study is dedicated to the quantitative
116 experimental validation of the developed nanocomposite-inspired broadband ultrasonic
117 sensor, capable of responding to acousto-ultrasonic waves (embracing AE and GUWs)
118 faithfully and accurately, either in a passive or active manner. Through constituent
119 optimization, the developed sensor is proven quantitatively responsive to GUWs up to 400
120 kHz. Lightweight and small, the sensor is demonstrated to surpass conventional ultrasonic

121 transducers such as PZT in some aspects, including the possibility to coat on a structure and
122 flexibility adapt to a curved structural surface, with reduced use of wiring and cabling. To
123 take a step further, the developed sensors are networked to form a dense sensor network,
124 well addressing an *in-situ* sensing philosophy and showing recommendable capability in
125 balancing “sensing cost” with “sensing effectiveness”.

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127 This paper is organized as follows. Section 2 briefly introduces the development, fabrication
128 and optimization of the nanocomposite sensor. In this section, the sensing mechanism of the
129 sensor in responding to acousto-ultrasonic waves, based on the tunneling effect in the
130 formed nanofiller conductive network, is explained and validated via material
131 morphological characterization tests. Section 3 concerns the conformance tests of the
132 developed sensor, in a broadband regime. For calibration, signals measured with the
133 nanocomposite sensor are compared with those obtained with conventional PZT wafers.
134 Upon conformance validation, the developed sensor is applied to damage identification
135 paradigms including impact localization (using low-frequency AE waves in a passive
136 manner) and quantitative damage evaluation (using high-frequency GUWs in an active way),
137 as detailed in Section 4, followed with concluding remarks in Section 5.

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139 **2. Sensor Fabrication and Characterization**

140 Targeting a broad range of response, the nanofiller and matrix of the nanocomposite sensor
141 were selected prudently according to twofold consideration: (i) the time- and
142 frequency-dependent viscoelastic properties of the candidate materials should be minute,
143 facilitating the sensor to be responsive to broadband acousto-ultrasonic waves from
144 low-frequency AE waves to high-frequency GUWs; and (ii) the gauge factor should be
145 sufficiently high, endowing the sensor with a capability to perceive GUWs of ultralow

146 magnitudes. A variety of nanofillers, including CB and CNT of various aspect ratios, has
147 been attempted, among which CB was selected as the nanofiller, owing to its lower aspect
148 ratio and therefore a higher specific surface and less amount of nanoparticle entanglement,
149 compared with other candidate CNT nanofillers. This attribute of CB is beneficial to the
150 formation of an even, stable and uniform conductive network in the sensor, to enhance the
151 sensitivity of sensor to broadband acousto-ultrasonic waves. On the other hand, PVDF was
152 chosen as the matrix, on the grounds that PVDF is a thermoplastic material with advantages
153 including easy-processing, extreme flexibility, low density, chemical inertness, thermal
154 stability, and good mechanical performance [29, 30, 47]. With a higher elastic modulus,
155 PVDF shows a faster response than rubber-based piezoresistive materials (rubber-based
156 matrix candidates often exhibit complex time- and frequency-dependent viscoelastic
157 properties, unwieldy to respond to dynamic loads).

158

159 Nanocomposite films were fabricated according to the process detailed in [28]. As shown
160 in Figure 1 (a), the film shows good resilience, highlighting its potential to adapt to a
161 curved structural surface. To further fabricate a nanocomposite film into a functional
162 sensor for acquisition of acousto-ultrasonic waves, the developed films were tailored to
163 individual rectangular flakes (10 mm × 8 mm, thickness remained at ~200 μm). Silver
164 conductive adhesive (D05001, Beijing Emerging Technology Co. Ltd., China) was pasted
165 on each flake, leading to two electrodes to which shielded cables were glued, as shown in
166 Figure 1 (b). It is noteworthy that the nanocomposite material between the two electrodes is
167 the sensor *per se*, in which the formed conductive network is responsive to GUWs.
168 Lightweight and flexible, each sensor, in a modality of thin film, can be produced via a
169 standard fabrication process, and deployed in a large quantity to configure a sensor
170 network for acquisition of acousto-ultrasonic waves. Repeating the above process, a series

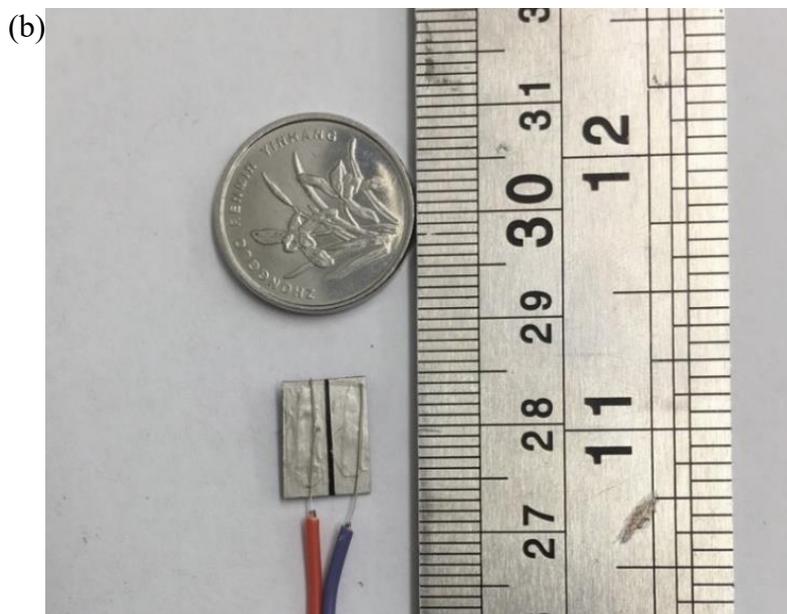
171 of nanocomposite sensor featuring various weight ratios of CB to PVDF, ranging from 5 to
172 30 wt%, were prepared in a comparative manner.

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178 **Figure 1** (a) A prepared nanocomposite film, showing good resilience; (b) a developed
179 sensor with silver-pasted electrodes and shielded cables

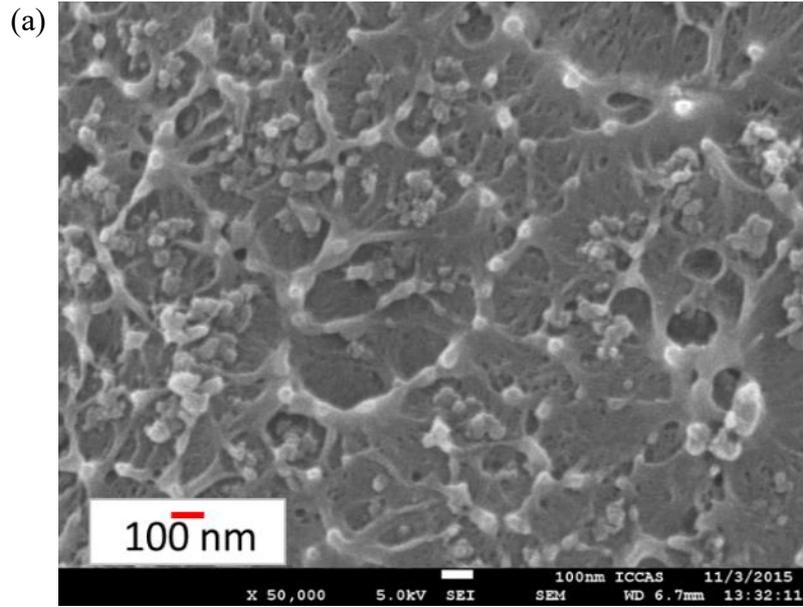
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181 To determine an optimal nano-structure of the sensor so as to enhance its sensing
182 sensitivity, a series of material characterization tests on the developed nanocomposites was
183 implemented. Electrical conductivity (σ) of the nanocomposite films with five
184 representative weight ratios of CB nanofiller (i.e., 5, 6.5, 8, 20 and 30 wt%, respectively)
185 was measured [28], which has affirmed that 6.5 wt% was the percolation threshold of the
186 CB conductive network. Percolation threshold represents a critical transition of the
187 nanocomposites from insulation to conduction, at which the nanocomposites exhibit the
188 highest sensitivity to external disturbance (e.g., the strain induced by acousto-ultrasonic
189 waves). Figure 2 (a) displays the scanning electron microscopy (SEM) (JSM-7500F, JEOL
190 Ltd) image of the nanocomposites (with ~6.5 wt% of CB nanofiller), revealing even and
191 uniform dispersion of nanoparticles in matrix PVDF.

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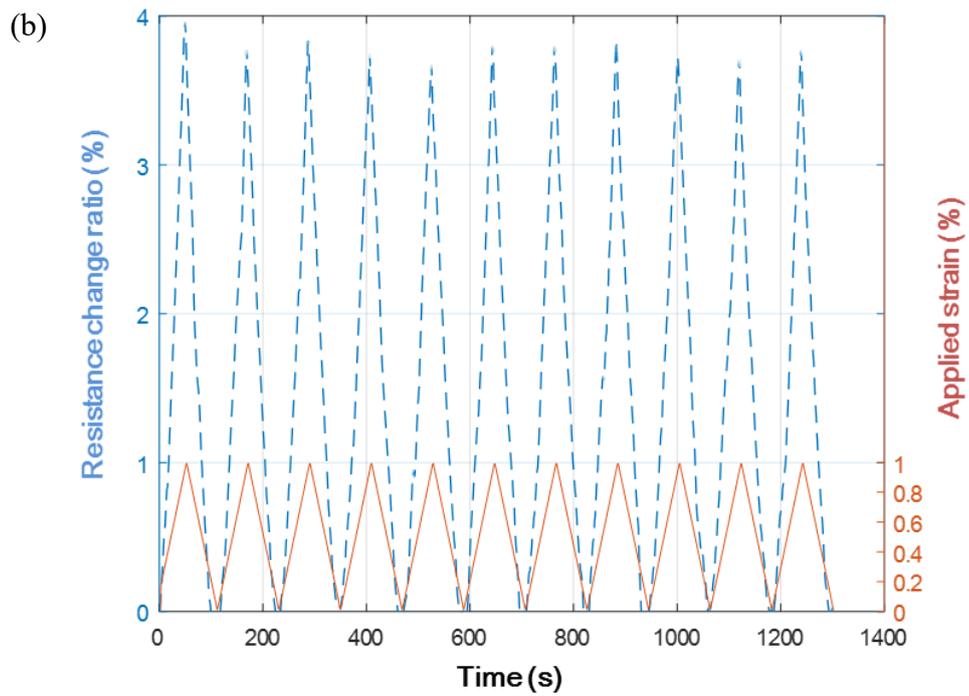
193 Upon determination of the percolation threshold, electro-mechanical properties of the
194 nanocomposites with 6.5 wt% of CB nanofillers were examined, using a semi-conductor
195 characterization system (4200-SCS, Keithley Instruments, Inc.) installed on a dynamic
196 mechanical testing platform (TA Q800, TA Instruments). σ of the nanocomposite films
197 was measured using a two-probe method when the films were subjected to a quasi-static
198 (1/120 Hz) cyclic tensile load until a maximum strain of 1% was reached (with a strain ramp
199 rate of 0.5%/min). Figure 2 (b) shows the obtained response of the nanocomposite films
200 against the cyclic load, to observe superb consistency between the response of the
201 nanocomposite films and the applied load, without observable hysteresis and distortion in
202 response waveform.

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207 **Figure 2** (a) SEM image of nanocomposites with ~6.5 wt% of CB nanofillers; and (b)

208 electro-mechanical response of prepared CB/PVDF nanocomposites (6.5 wt% of CB

209 nanofillers) subjected to a quasi-static load (1/120 Hz)

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212 To extend the above examination from a quasi-static (1/120 Hz) frequency to higher
213 frequencies, a developed nanocomposite film sensor (6.5 wt% of CB nanofillers) was
214 surface-bonded on a glass fiber/epoxy composite beam (290 mm long, 38 mm wide and 2
215 mm thick), as shown in Figure 3 (a). The beam was excited 270 mm from the clamped end
216 via an electro-mechanical shaker (B&K[®] 4809) with a sinusoidal signal of 2000 Hz. For the
217 purpose of calibration and comparison, a commercially available metal-foil strain gauge
218 (with a gauge factor of ~2) was collocated aside the nanocomposite sensor, both having the
219 same distance of 200 mm to the excitation point.

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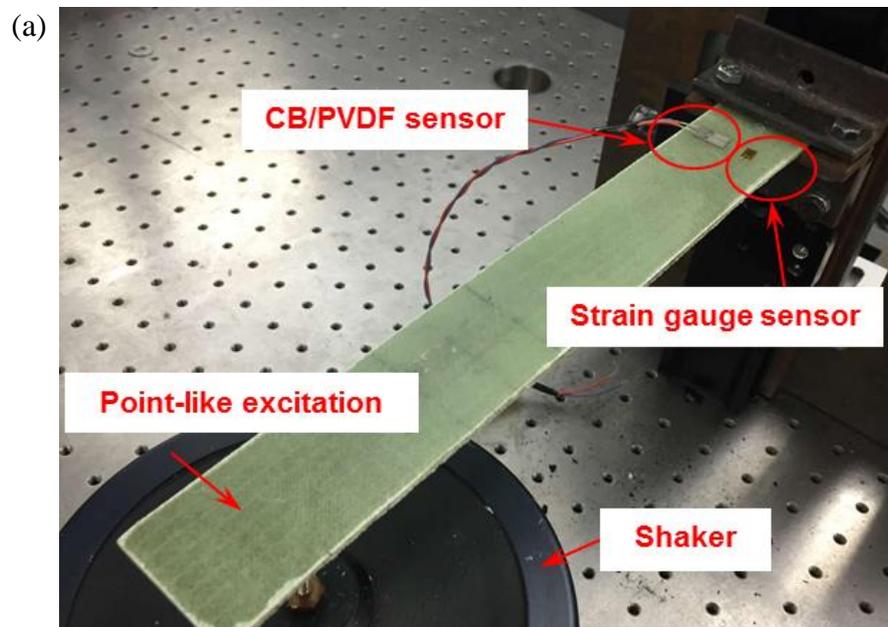
221 The magnitude of an acousto-ultrasonic wave can be faint, particularly when it is in a
222 frequency range of kilo- or mega-Hertz – that is the case for GUWs used for
223 acousto-ultrasonics-based SHM, in which the magnitude of the GUW-induced strain is
224 usually of several micro-strains only. With such a low degree of magnitude, a captured
225 acousto-ultrasonic wave signal can be naturally prone to contamination from environment
226 noise and measurement uncertainties, leading to a low signal-to-noise ratio. To this end, a
227 self-developed signal amplification module was configured, to be used in conjunction with
228 the sensor. The signal amplification module mainly integrates a Wheatstone bridge with
229 adjustable resistors compatible with the electrical resistance of the nanocomposite sensor, an
230 electronic amplifier circuit, a series of high-pass and low-pass filters and a signal conversion
231 circuit for converting measured piezoresistivity to electrical signals.

232

233 Using an oscilloscope (Agilent[®] DSO9064A), acousto-ultrasonic wave signals acquired
234 with both the nanocomposite sensor and strain gauge were registered, with an example
235 displayed in Figure 3 (b). Both signals show high consistency, as well as good reversibility
236 and repeatability of measurement. Observation from this electro-mechanical test argues that

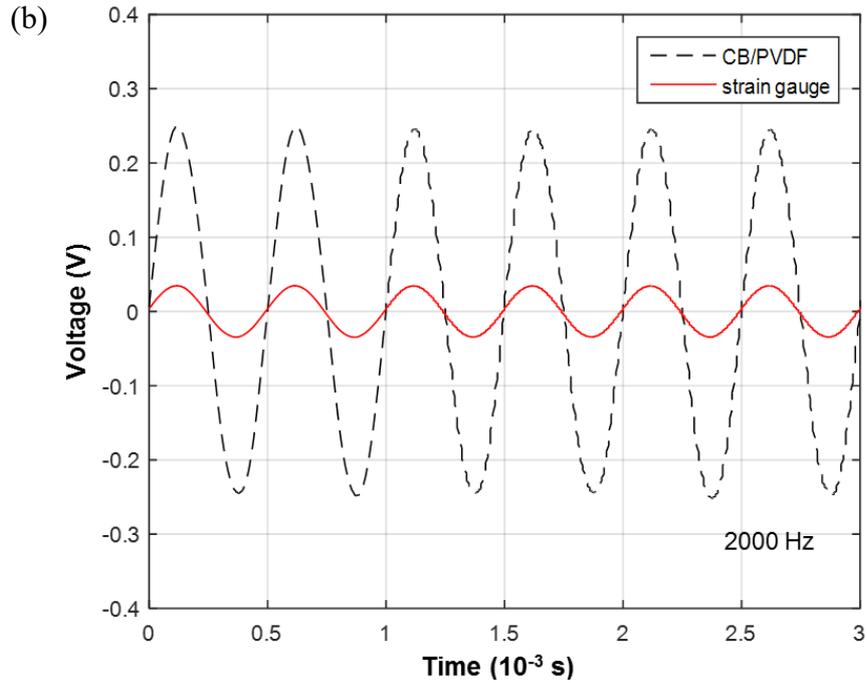
237 the fabricated sensor made with CB/PVDF nanocomposites at its percolation threshold (~6.5
238 wt% of CB nanofillers) is responsive to a dynamic excitation up to 2000 Hz, without
239 observable hysteresis and deviation in response waveform and with a greater gauge factor
240 (~10) five times larger than conventional strain gauges (~2). It is noteworthy that the gauge
241 factor increases as the sensing frequency (~4 in the quasi-static cyclic loading of 1/120 Hz
242 and ~10 in the vibration test of 2000 Hz), which facilitates the application of the developed
243 CB/PVDF sensor for perception of acousto-ultrasonic waves.

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249 **Figure 3** (a) Experiment set-up for acquisition of vibration signal of a glass fiber/epoxy
250 composite beam using the prepared nanocomposite sensor (6.5 wt% of CB nanofillers); and
251 (b) comparison of signals captured with nanocomposite sensor and metal-foil strain gauge
252 (at 2000 Hz)

253

254 To achieve insight into the sensing mechanism of the developed nanocomposite sensor in
255 responding to broadband acousto-ultrasonic waves, an X-ray diffraction (XRD) test [28]
256 was accomplished, to confirm that the prepared CB/PVDF nanocomposites feature a pattern
257 of α -crystal with a non-polar crystal structure. This suggests that the piezoresistivity
258 manifested by the developed sensor is from the change in nanofiller-formed conductive
259 network caused when an acousto-ultrasonic wave traverses the sensor, rather than from the
260 piezoelectricity effect of PVDF matrix itself. This is because only a polar crystal (such as a
261 β -crystal) can lead to piezoelectricity (as interpreted elsewhere [35]). Based on XRD results,
262 conclusion can be drawn that the predominant sensing mechanism of the developed
263 nanocomposite sensor is the tunneling effect [31-33] present in the conductive network

264 formed by CB nanoparticles (as well as their aggregations) when the sensor is exposed to an
265 acousto-ultrasonic wave. In brief, the tunneling effect, at nanoscale, occurs when
266 neighboring nanoparticles are in a close proximity (normally several nanometers) but not in
267 a direct contact. When the sensor (at the determined percolation threshold) is subjected to an
268 acousto-ultrasonic wave, the distances among adjacent nanoparticles alter, leading to the
269 tunneling effect of charged carriers. As a direct consequence of the tunneling effect, the
270 conductivity of the nanofiller network and consequently the electrical resistance measured
271 by the sensor varies – that is the primary sensing mechanism of the sensor in responding to
272 broadband acousto-ultrasonic waves.

273

274 **3. Sensor Calibration for Acquisition of Acousto-ultrasonic Waves**

275 Upon determination of an optimal nano-structure, the developed CB/PVDF nanocomposite
276 film sensor was calibrated for *in-situ* perception of acousto-ultrasonic waves ranging from
277 low-velocity-impact-induced low-frequency AE waves to active high-frequency GUWs.

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279 **3.1. Experiment Set-up**

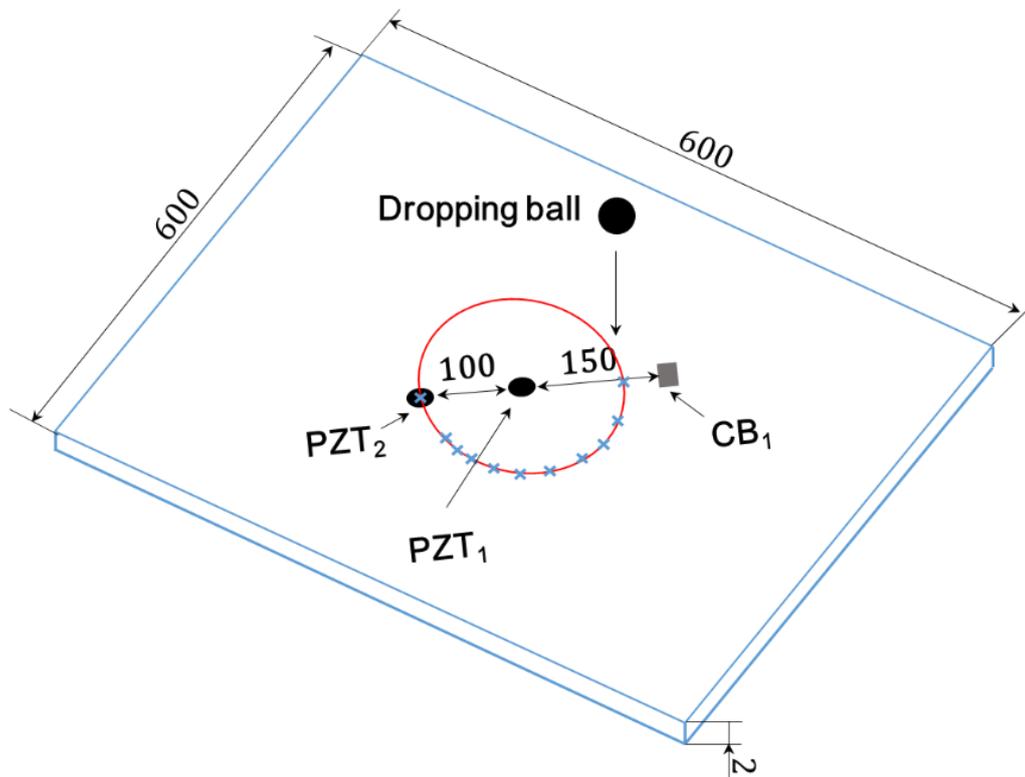
280 A glass fiber/epoxy composite panel (600 mm × 600 mm × 2 mm) was prepared with its
281 four edges clamped, to which a fabricated nanocomposite sensor (denoted by CB₁
282 hereinafter) and two commercial PZT wafers (Physik Instrumente Co., Ltd., PIC151; 9 mm
283 in diameter and 0.5 mm in thickness; denoted by PZT₁ and PZT₂) were surface-mounted
284 with an instant glue, shown schematically in Figure 4. PZT₁, centralized on the panel with
285 a distance of 150 mm to the nanocomposite sensor, was used to capture low-frequency AE
286 signals in a passive drop test (Section 3.2) or to generate high-frequency GUWs in an
287 active GUW test (Section 3.3), while PZT₂, 100 mm from PZT₁, functioned as a sensor to
288 acquire active GUWs generated by PZT₁ (in Section 3.3).

289 Due to different sensing philosophies of these two types of sensor (piezoelectric effect for
290 PZT wafer, while tunneling effect-induced piezoresistive effect for nanocomposite sensor
291 as interpreted in Section 2), different signal acquisition methods were adopted: PZT wafers
292 were instrumented with an oscilloscope (Agilent® DSO9064A), while CB/PVDF with the
293 self-developed test system as detailed in Section 2.

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298 **Figure 4** Schematic of experimental set-up for *in-situ* perception of acousto-ultrasonic

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waves (unit: mm)

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304 3.2. Acquisition of Low-frequency AE Signals

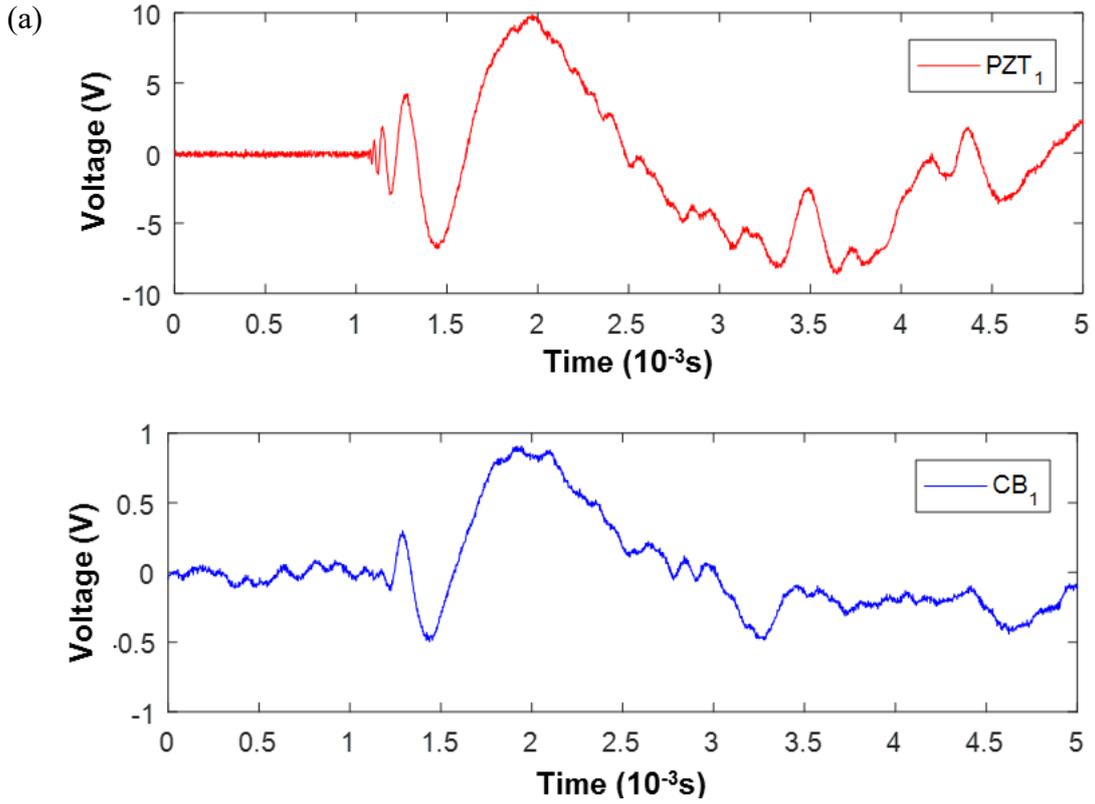
305 A drop test was performed using the experimental set-up in Figure 4, in which a steel ball
306 (8 mm in diameter) impinged on the panel from a height of 300 mm to introduce an impact
307 energy of $\sim 6.3 \times 10^{-3}$ J to the panel. The generated AE waves were then captured with both
308 sensors (CB_1 and PZT_1) at a sampling rate of 1 MHz. The drop test was repeated,
309 introducing impact energy to the panel at different spots along the periphery of a circle
310 with a radius of 100 mm and PZT_1 being the center (symbol “×” in Figure 4), this leading
311 to different distances from the impact spot to the nanocomposite sensor, varying from 50 to
312 250 mm.

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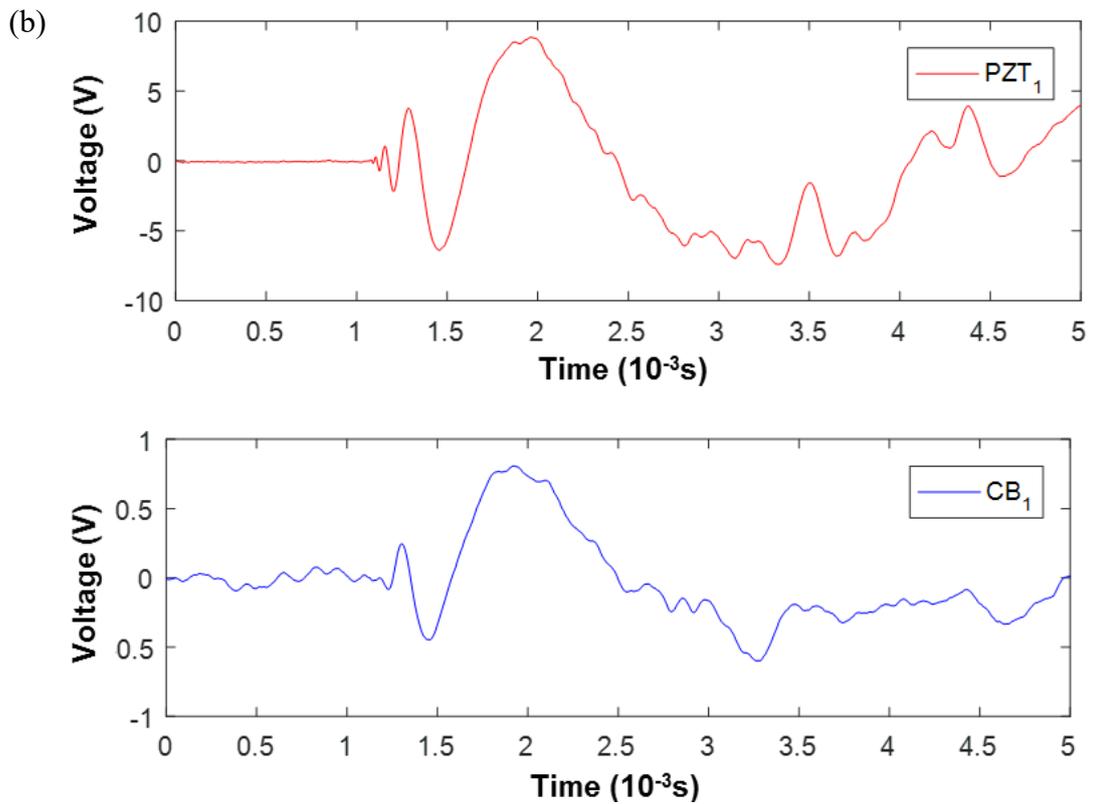
314 A set of typical AE signals acquired with PZT_1 and CB_1 , at a respective distance of 100
315 mm and 90 mm to the impact spot, is shown in Figure 5 (a). To mitigate measurement
316 noise, all raw signals were processed with a first-order Butterworth low-pass filter with a
317 cutoff frequency of 10 kHz, and the accordingly processed signals are presented in Figure
318 5 (b), to observe a good coincidence in performance between the nanocomposite sensor
319 and the PZT wafer. The zeroth order anti-symmetric mode (A_0) is observed to be the
320 dominant propagating wave mode in the low-velocity-impact-induced AE signals. Further,
321 Figure 6 compares the spectra of the two signals in Figure 5, affirming that the
322 nanocomposite sensor responds to a signal of a frequency as high as 5 kHz, with
323 comparable performance with commercial PZT-type sensors.

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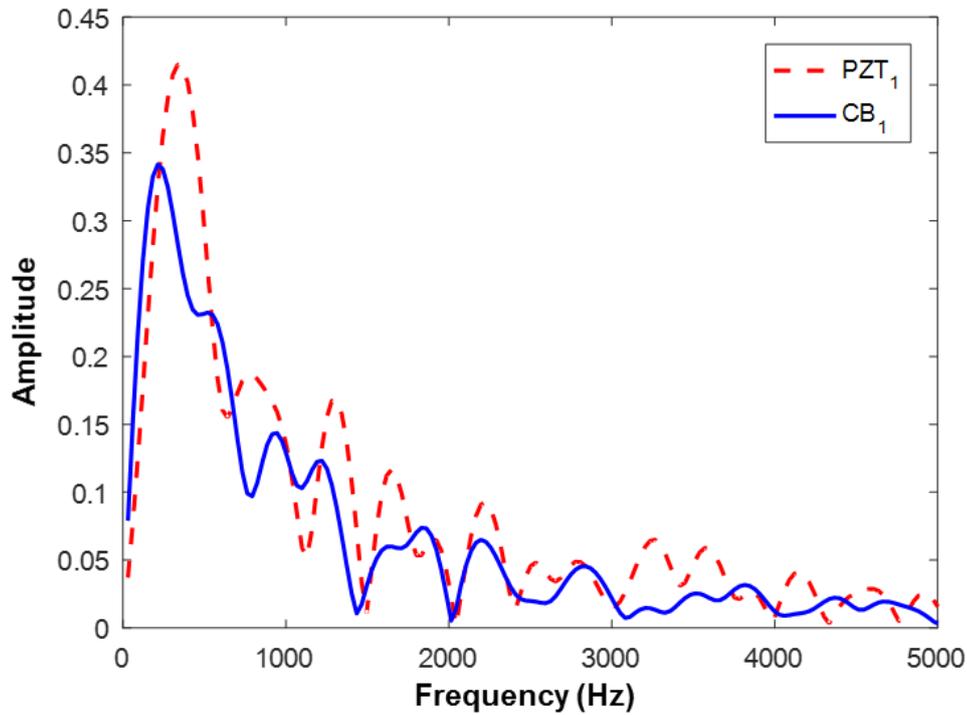
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328 **Figure 5** (a) Raw and (b) noise-filtered AE wave signals acquired with PZT₁ and CB₁ at a
329 respective distance of 100 mm and 90 mm to the impact spot

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331 **Figure 6** Comparison of spectra of signals in Figure 5 acquired with PZT₁ and CB₁

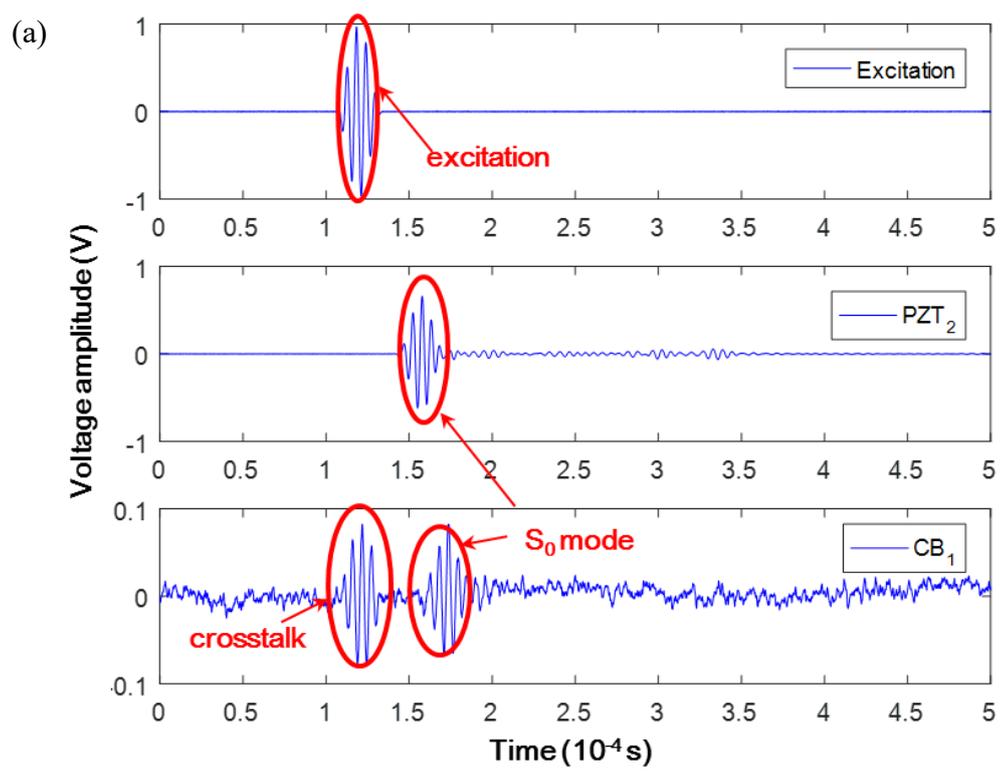
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333 3.3. Acquisition of High-frequency GUV Signals

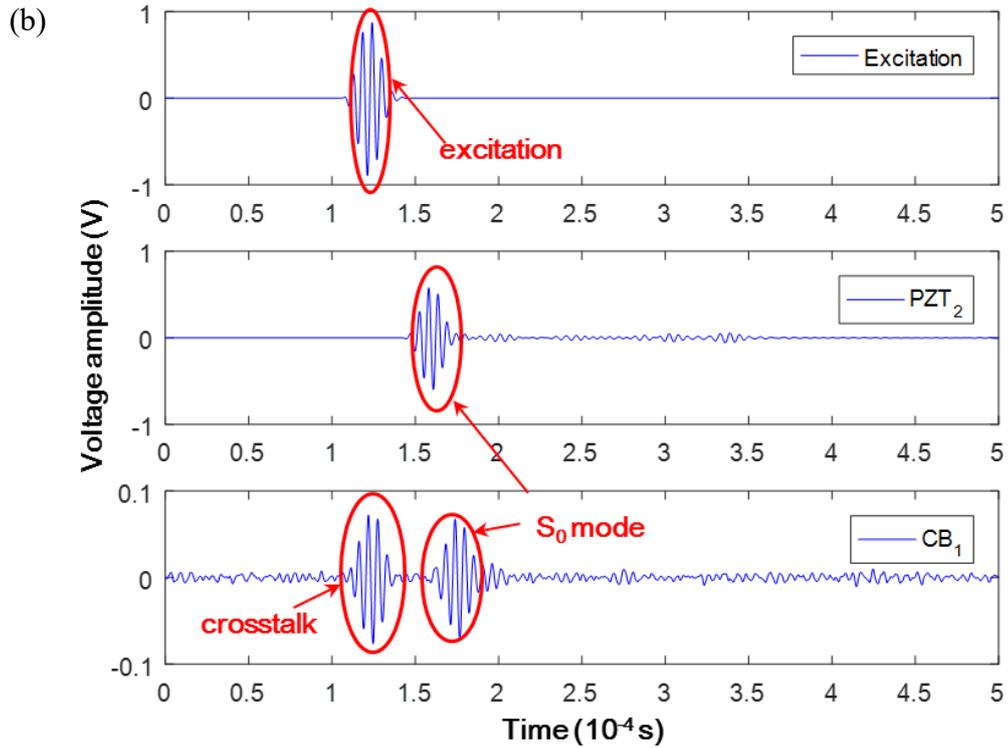
334 GUVs in an ultrasonic regime feature extremely low strains (of the order of micro-strain)
335 whereas ultrafast dynamic change (>20 kHz). The acquisition of high-frequency GUV
336 signals using the developed nanocomposite sensor was validated, with the panel in Figure 4.
337 The GUV generation and acquisition system includes a waveform generator (NI®
338 PXI-5412 integrated in the NI® PXIE-1071 4-slot chassis), a linear power amplifier
339 (Ciprian® US-TXP-3), and the self-developed signal amplification and acquisition module
340 as detailed in Section 2. In the test, the wave generator output five-cycle
341 Hanning-windowed sinusoidal tone-bursts at central frequencies from 50 kHz to 400 kHz.
342 Upon amplification to 400 V_{p-p} to the linear power amplifier, the tone-bursts excited PZT₁
343 to produce GUVs propagating in the panel, which were then acquired with CB₁ and PZT₂,
344 respectively, at a sampling rate of 20 MHz. Notably, in order to eliminate possible

345 interference from the PZT on the nanocomposite sensor, different wave propagation
 346 distances were set deliberately to two types of sensor: 100 mm from PZT₁ to PZT₂, and
 347 150 mm from PZT₁ to CB₁.
 348
 349 At a representative frequency of 200 kHz, the excitation signal and accordingly acquired
 350 GUV signals with CB₁ and PZT₂ are shown in Figure 7. The raw signal acquired with CB₁,
 351 Figure 7 (a), is noted to embrace the zeroth order symmetric Lamb wave mode (S₀) – the
 352 first-arrival wave component, and a high-voltage (400 V_{p-p}) crosstalk induced by the
 353 linear power amplifier. Owing to distinct sensing mechanisms – the piezoelectric effect for
 354 PZT and piezoresistive effect for CB/PVDF – the signal acquired with PZT shows a higher
 355 magnitude and higher signal-to-noise ratio. Upon applied with a Butterworth filter (Figure 7
 356 (b)) and a Hilbert transform (Figure 7 (c)), both signals become more explicit and consistent,
 357 to observe good and quantitative agreement in between, in terms of the waveform and arrival
 358 moment of the concerned wave mode. The filtered signals were then used for damage
 359 detection.

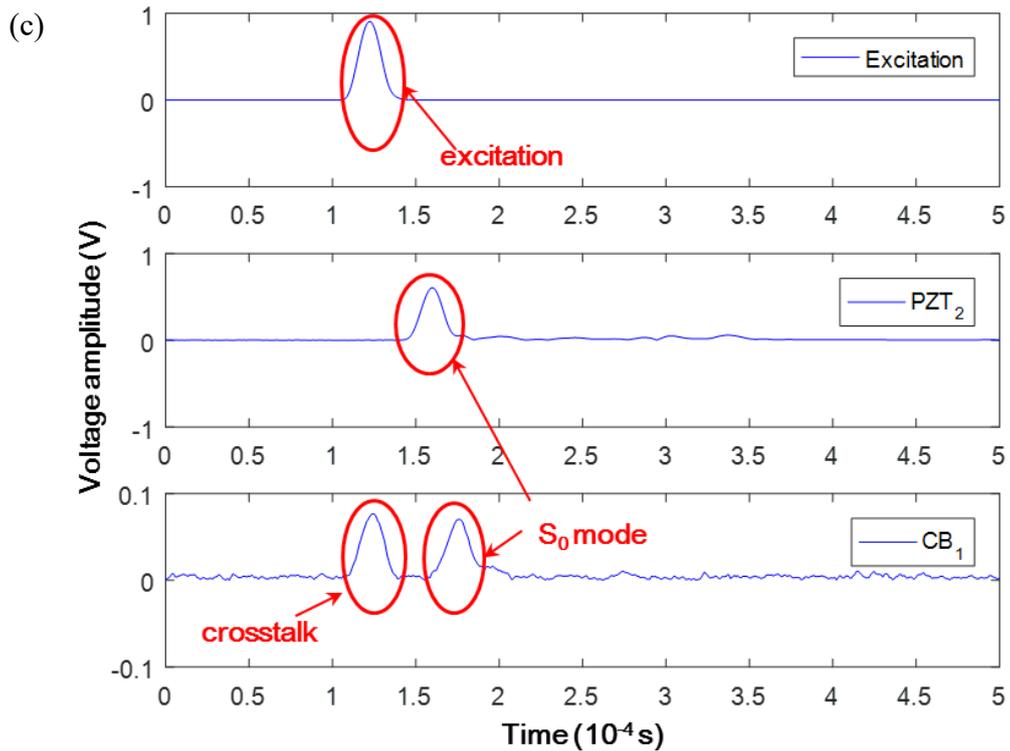
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363 **Figure 7** (a) Raw GUW signals captured with PZT₂ and CB₁ at 200 kHz; (b) filtered signals
364 using Butterworth filter; and (c) processed signals using Hilbert transform (Note: the arrival
365 time of S₀ mode in PZT₂- and CB₁-captured signals is different because of the different
366 distances from PZT₁ to PZT₂ and to CB₁)

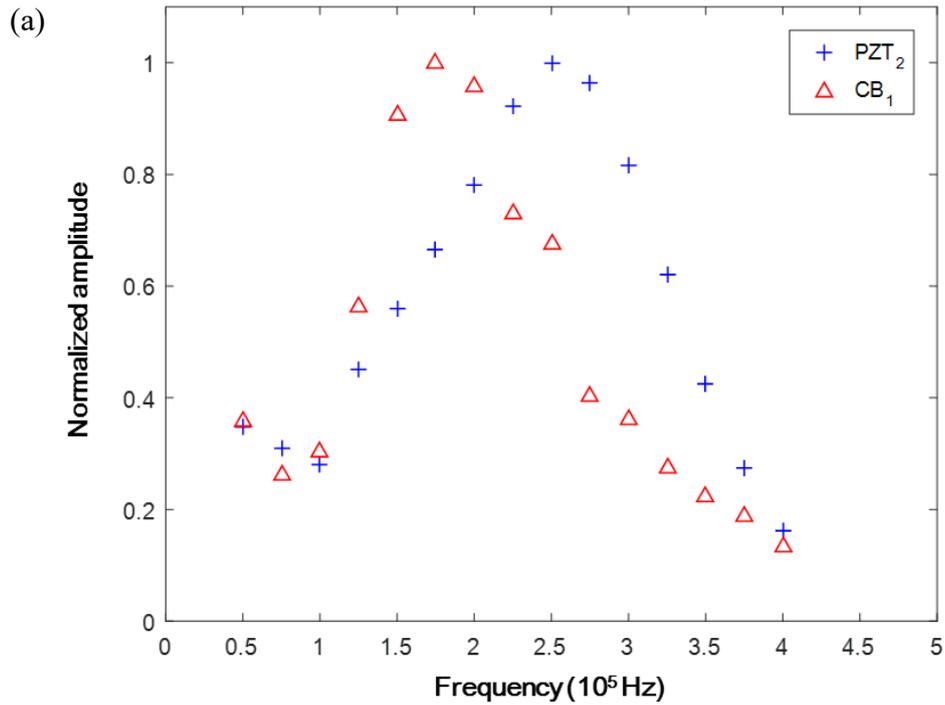
367 To extend the above discussion (at 200 kHz), a series of tone-bursts from 50 kHz to 400
368 kHz was excited via PZT₁, and Figure 8 (a) displays the normalized magnitudes of the
369 signals acquired by PZT₂ and CB₁, respectively. The nanocomposite sensor is observed to
370 reach a response peak at 175 kHz within the discussed frequency range, compared with
371 PZT at 250 kHz – a property known as “mode tuning” (i.e., the magnitude of the sensor
372 response varies subjected to excitation frequency) [11, 48]. The difference in mode tuning
373 between two sensors (175 kHz for nanocomposites while 250 kHz for PZT) could be
374 attributed to the different sensing mechanisms of nanocomposite sensor and PZT.

375

376 With G UW signals in Figure 8 (a), Figure 8 (b) further compares the ascertained
377 propagation velocities of S₀ and A₀ at various excitation frequencies, to note a coincident
378 performance of two types of sensor in responding to various G UW mode in a wide
379 frequency range as high as 400 kHz, with a weak magnitude as low as ~ 10 micro-strain.
380 This has testified the superior capacity of the developed nanocomposite sensor as a
381 broadband ultrasonic sensor.

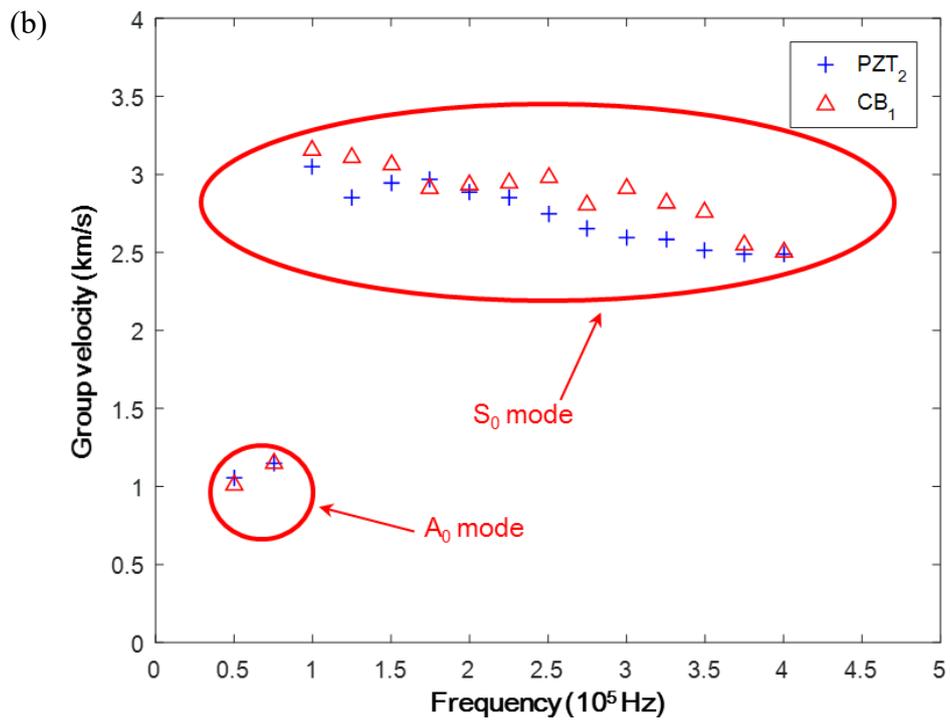
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387 **Figure 8** Variation in response of prepared CB/PVDF nanocomposite sensor (6.5 wt% of CB

388 nanofillers) to excitation, compared with PZT: (a) normalized magnitude vs. excitation

389 frequency; (b) group velocity vs. excitation frequency

390 **4. Applications to Passive and Active Damage Characterization**

391 Upon material characterization, optimization, performance calibration and conformance
392 tests, the developed CB/PVDF nanocomposite film sensor was applied to two paradigms of
393 damage characterization – passive impact localization and active damage identification. In
394 both paradigms, the sensor performed as a broadband ultrasonic sensor for *in-situ*
395 acquisition of acousto-ultrasonic waves. In addition, being lightweight and flexible,
396 individual film sensors were networked in a dense fashion, to render rich information for
397 depicting damage with desirable redundancy and hence enhanced reliability of signal
398 acquisition, outperforming traditional piezoelectric counterpart in terms of adaptability to
399 complex structure and information redundancy. With the sensor network, damage was
400 evaluated quantitatively with parameters including presence, location and severity.

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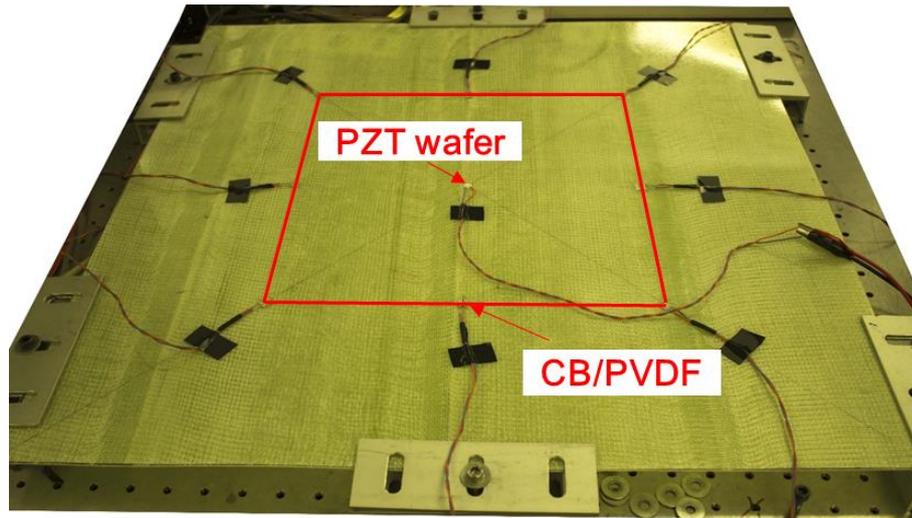
402 **4.1. Passive Paradigm: Impact Localization**

403 Eight nanocomposite sensors (denoted by CB₁~CB₈) were surface-glued on an
404 all-boundary-clamped fiber-epoxy composite plate, see Figure 9, to form a sensing network
405 with an inspection coverage of approximately 300 mm × 300 mm. A drop test with the
406 same configuration used earlier (Section 3.1), including the same degree of impact energy,
407 was recalled. In the test, a steel ball impinged on the plate at eight locations, as marked
408 with symbol “cross” (×) in Figure 9 (b), each of which had the same distance of 100 mm to
409 a PZT wafer centralized on the plate (denoted by PZT₁). The procedure and configuration
410 of signal acquisition were remained the same as those used in Section 3.1.

411

412

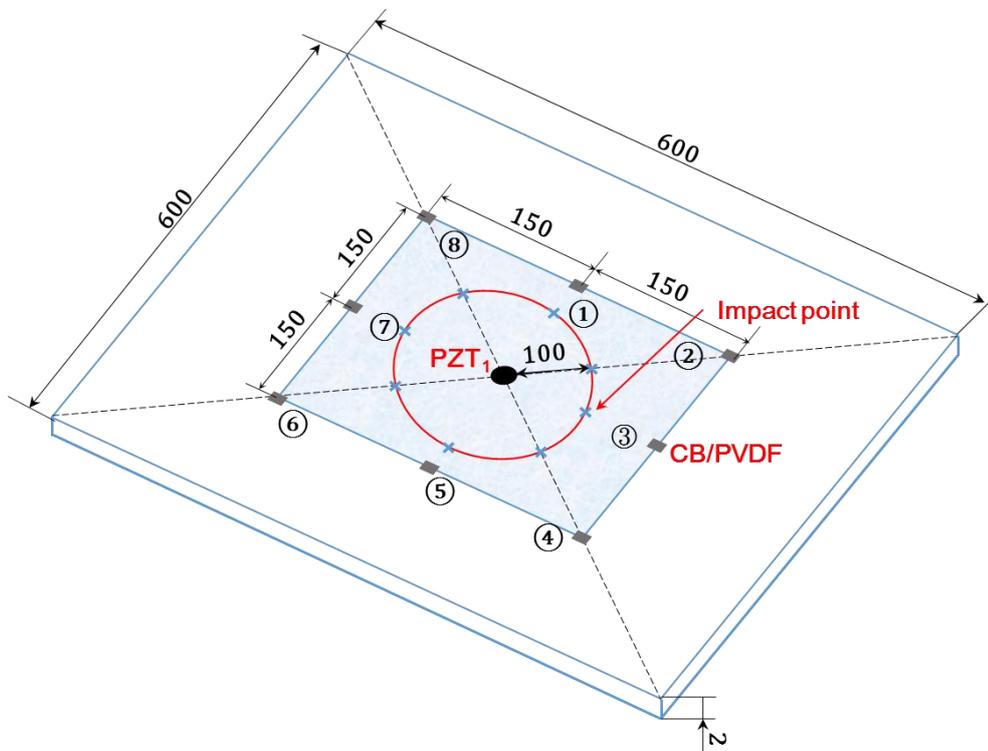
(a)



413

414

(b)



415

416 **Figure 9** (a) Photographed and (b) schematic description of experimental set-up (with a
417 surface-glued sensor network formed by nanocomposite film sensors), for passive impact
418 localization and active damage identification (unit: mm)

419

420

421 With obtained AE wave signals, a *delay-and-sum* imaging algorithm [49] was recalled to
 422 locate impact spots in eight drop tests. In the algorithm, the first-arrival wave component
 423 captured with the i^{th} sensor, located at (x_i, y_i) , in the sensor network (denoted by CB_i) is
 424 expressed as

$$425 \quad t_i = t_0 + \Delta t_i, \quad (1)$$

426 where t_0 is the moment when the steel ball impacts the plate, and Δt_i the subsequent
 427 traveling time for A_0 wave mode in the signal from the impact spot to CB_i (it has been
 428 earlier demonstrated in Section 3.2 that A_0 mode dominates a low-velocity-impact-induced
 429 AE signal).

430

431 Taking into account another sensor in the sensor network, say CB_j at (x_j, y_j) , and assuming
 432 the impact spot is at (x, y) within the inspection coverage, the time difference ($\Delta t_{ij}(x, y)$)
 433 in the arrival time of A_0 captured by CB_i and CB_j yields

$$434 \quad \begin{aligned} \Delta t_{ij}(x, y) &= t_i - t_j = (t_0 + \Delta t_i) - (t_0 + \Delta t_j) \\ &= \frac{\sqrt{(x - x_i)^2 + (y - y_i)^2} - \sqrt{(x - x_j)^2 + (y - y_j)^2}}{v_{plate}}. \end{aligned} \quad (2)$$

435 In Eq. (2), variables are distinguished by subscripts i and j for two individual sensors.
 436 v_{plate} represents the velocity of A_0 mode.

437

438 With the sensor pair formed by CB_i and CB_j , a two-dimensional gray-scale image can be
 439 synthesized using the *delay-and-sum* algorithm, with its pixel value ($\xi_{ij}(x, y)$) defined as

$$440 \quad \xi_{ij}(x, y) = \max(E_i + E_j(\Delta t_{ij}(x, y))), \quad (3)$$

441 where E is the energy packet of A_0 mode obtained using a wavelet transform. With a
 442 compensation for the time delay of $\Delta t_{ij}(x, y)$ in the arrival time of A_0 mode in the signal
 443 captured by CB_j , with regard to that in the signal captured by CB_i , the max operator in Eq.

444 (3) defines the peak value of the summation of two time-series signals acquired with CB_i
 445 and CB_j , which is then linked to the probability of existence of impact spot within the
 446 inspection coverage – the perceptions as to the impact spot from the perspective of the sensor
 447 pair CB_i and CB_j . According to Eqs. (2) and (3), all the pixels in the image which feature the
 448 same time delay will form a set of hyperbola, with CB_i and CB_j being its two foci. The
 449 hyperbola suggests all the possible locations of impact within the inspection coverage. In
 450 principle, those pixels with greater value of $\xi_{ij}(x, y)$ have higher degrees of probability of
 451 impact spot therein, and vice versa.

452

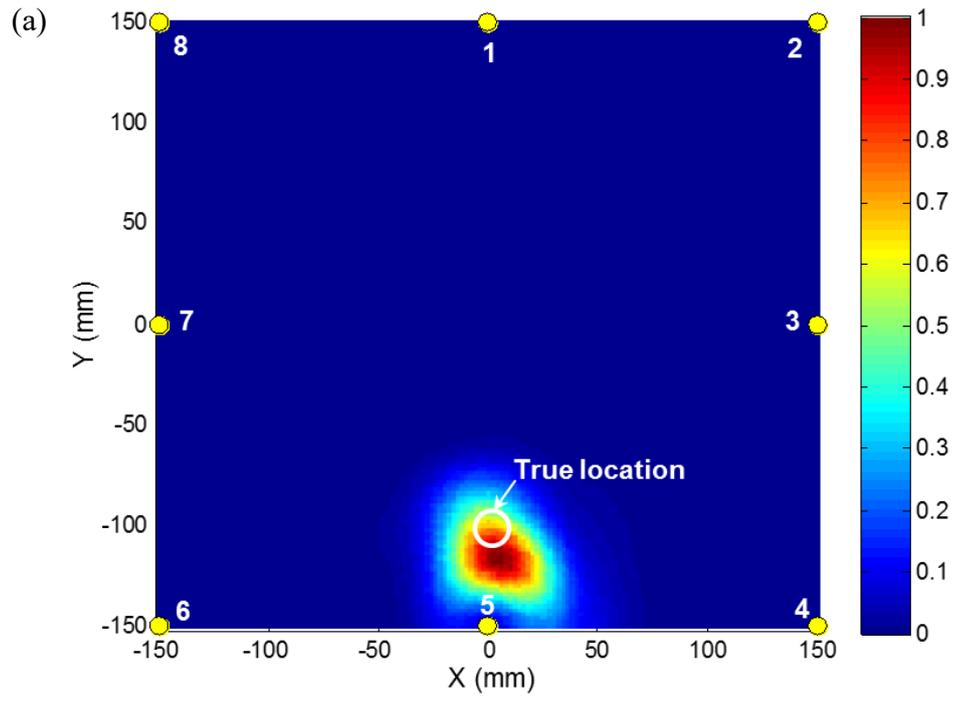
453 Aggregating images constructed by all available sensor pairs in the sensor network through
 454 appropriate image fusion algorithms, a superimposed image can be hypothesized – a
 455 collective consensus as to the impact spot from the entire sensor network. Without loss of
 456 generality, taking the AE signal acquired with CB_1 as reference, one has, for the current
 457 sensor network involving eight nanocomposite sensors,

$$458 \quad \xi_{sum}(x, y) = \max\left(\sum_{j=2}^8 (E_1 + E_j(\Delta t_{1j}(x, y)))\right). \quad (4)$$

459 As representative results, Figure 10 (a) and (b) exhibit the identified impact spots in two out
 460 of eight drop tests, in which a nonlinear normalization process was applied for enhanced
 461 focusing. Figure 10 (c) shows the combined results for all the eight tests. A high degree of
 462 coincidence between identified impact spots and reality can be noted, affirming the
 463 performance of the developed nanocomposite sensor in passive AE-based damage
 464 localization.

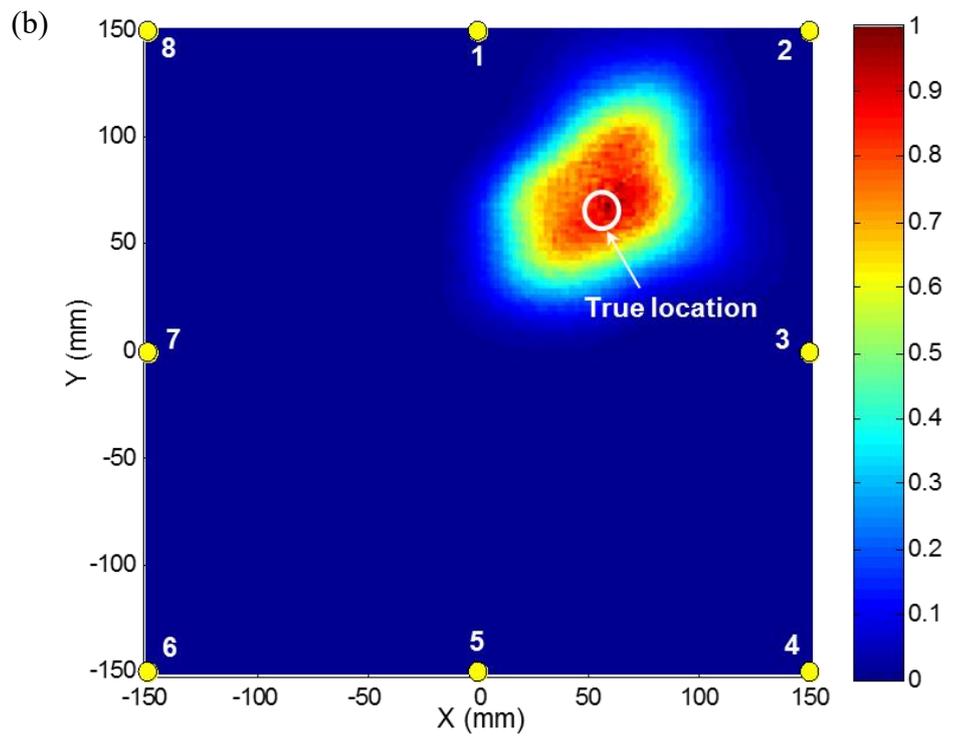
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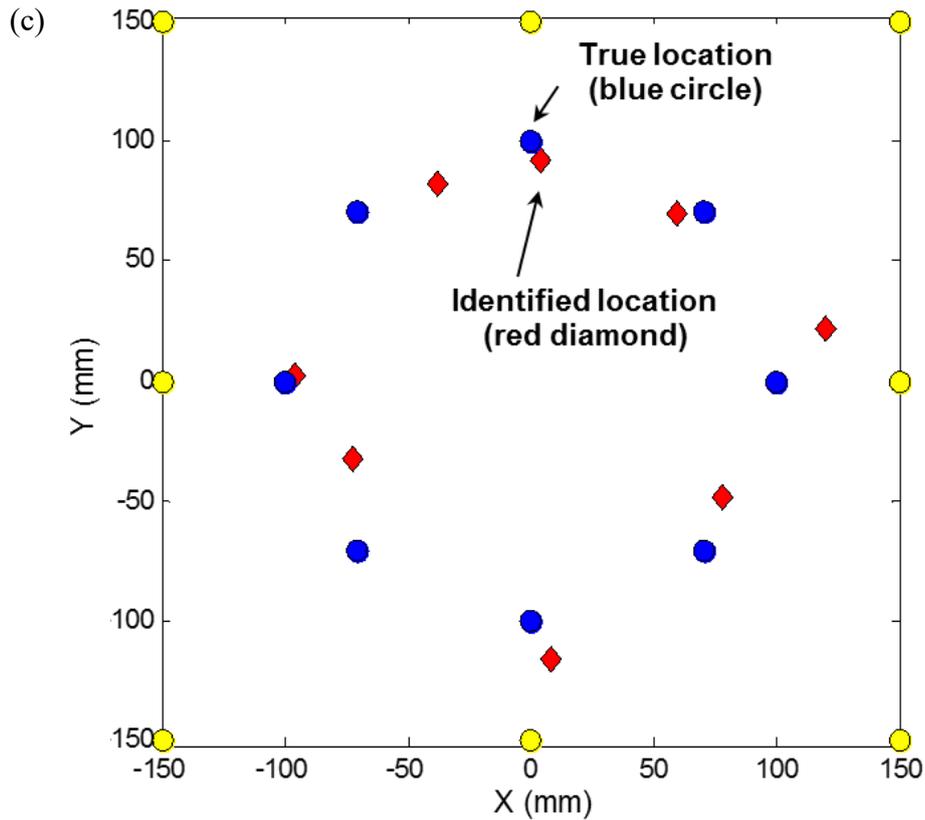


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470

471 **Figure 10** Comparison between identified and real impact spots using impact-induced
472 passive AE signals captured with the nanocomposite sensors, when the real spot is at (a) (0,
473 -100); (b) (71, 71); and (c) combined results for eight drop tests (unit mm)

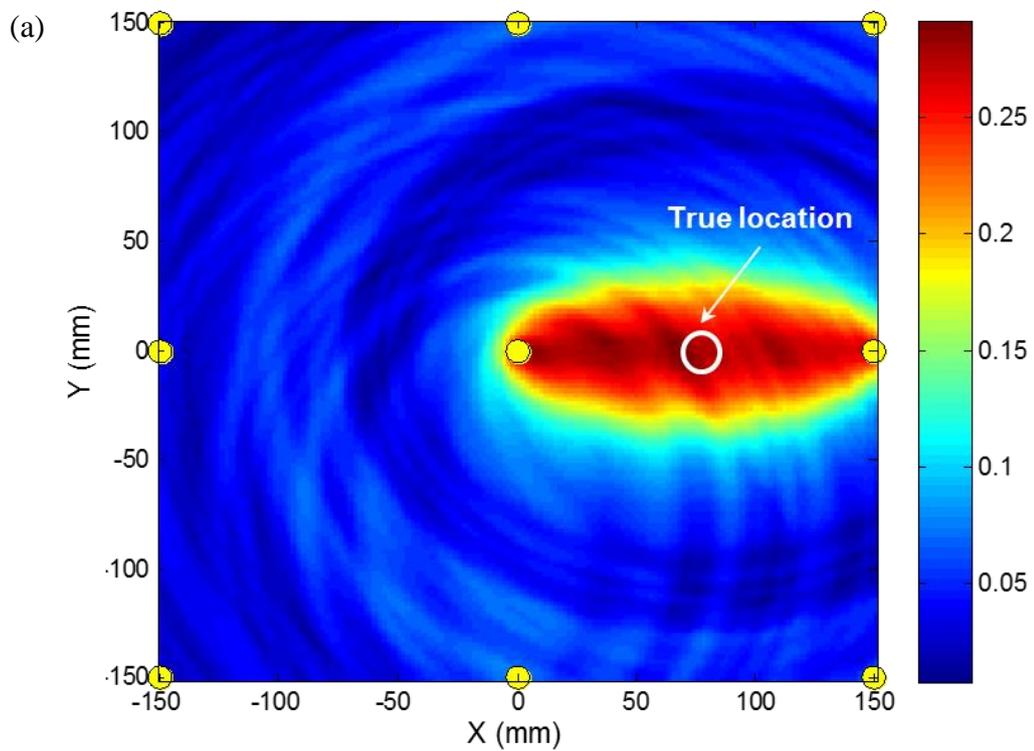
474

475 **4.2. Active Paradigm: Quantitative Characterization of Mock-up Damage**

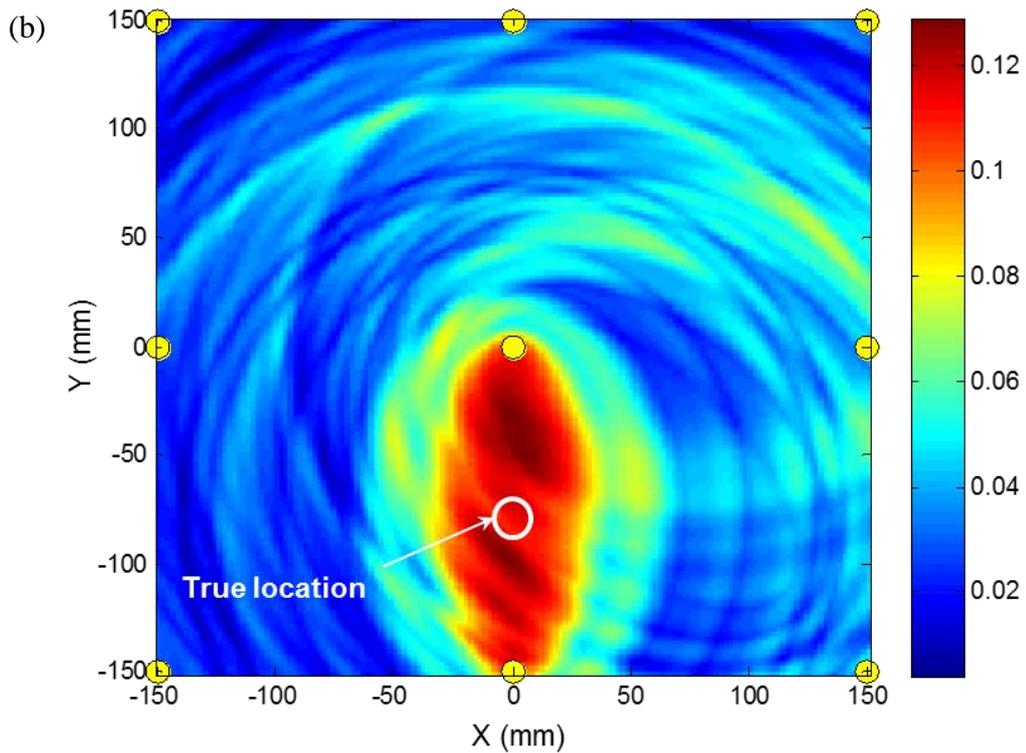
476 Using the same glass fiber-epoxy composite plate in Section 4.1, PZT₁, as a wave
477 generator, produced a five-cycle Hanning-windowed sinusoidal tone-burst at a central
478 frequency of 200 kHz with a magnitude of 400 Vp-p. The selection of 200 kHz lies in the
479 fact that at this frequency, S₀ mode becomes predominant in GUWs, according to the
480 “mode tuning” effect (Section 3.3). It is noteworthy that different sensing mechanisms
481 between the developed nanocomposite sensor and the PZT sensor decide the different
482 “mode tuning” effect, through which 200 kHz was selected to ensure a high signal-to-noise

483 ratio. The propagation velocity of S_0 mode was ascertained in Figure 8. A mock-up damage
484 (simulated using an adhesive tape measuring $20\text{ mm} \times 20\text{ mm} \times 3\text{ mm}$) was introduced to
485 the composite plate. The generated GUWs were acquired with the eight CB/PVDF
486 nanocomposite sensors at a sampling rate of 20 MHz. The *delay-and-sum*-based imaging
487 algorithm, as elaborated in Section 4.1, was applied. As representative results, Figure 11
488 shows the identification results for two scenarios when the mock-up damage was at (75
489 mm, 0 mm) and (0 mm, -75 mm), respectively. Though not pinpointing the damage
490 quantitatively and accurately (due to the use of only a few sensors), the identification
491 results are still able to point out the region where the mock-up damage exists. This has
492 substantiated a wide application domain of the developed nanocomposite sensor from
493 passive AE-based impact localization to active GUWs-based quantitative damage
494 characterization.

495
496



497



498

499 **Figure 11** Comparison between identified and real damage using actively generated GUWs
500 captured with the nanocomposite sensors, when the real damage is at (a) (75, 0); and (b) (0,
501 -75) (unit mm)

502

503 **5. Concluding Remarks**

504 In this study, the previously developed nanocomposite sensors, with the tunneling effect
505 being the dominant sensing mechanism, were networked for implementation of
506 acousto-ultrasonics-based damage characterization. Through the optimization of
507 nano-structure and nanofiller constituent (~6.5 wt%) of the sensor, the sensing performance
508 of the nanocomposite-inspired *in-situ* broadband ultrasonic sensor made of CB/PVDF was
509 validated, and experimentally evaluated quantitatively to perceive broadband response with
510 frequencies up to 400 kHz: from low-frequency vibration (2000 Hz), through
511 impact-induced AE waves (<5 kHz), to GUWs (up to 400 kHz), with high fidelity, ultrafast

512 response and high sensitivity. The sensor presents a much greater gauge factor (~ 10) under
513 vibration of 2000 Hz compared with conventional metal-foil strain gauge (~ 2), and greater
514 flexibility in accommodating curved structural surfaces. Lightweight, flexible, coatable to a
515 structure and deployable in a large quantity to form a dense sensor network for *in-situ*
516 acquisition of acousto-ultrasonic waves in either a passive or active manner, this new type
517 of broadband ultrasonic sensor has unfolded its application prospect towards
518 acousto-ultrasonics-based SHM.

519

520 **Acknowledgements**

521 This project is supported by the Hong Kong Research Grants Council via General Research
522 Fund (No. 15214414 and No. 15201416).

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