Characterization of Tissue Response to Impact Loads Delivered Using a Hand-Held Instrument for Studying Articular Cartilage Injury

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Abstract

Objective: The objective of this study was to fully characterize the mechanics of an *in vivo* impactor and correlate the mechanics with superficial cracking of articular surfaces. *Design:* A spring-loaded impactor was used to apply energy-controlled impacts to the articular surfaces of neonatal bovine cartilage. The simultaneous use of a load cell and displacement sensor provided measurements of stress, stress rate, strain, strain rate, and strain energy density. Application of India ink after impact was used to correlate the mechanical inputs during impact with the resulting severity of tissue damage. Additionally, a signal processing method to deconvolve inertial stresses from impact stresses was developed and validated. *Results:* Impact models fit the data well (root mean square error average ~0.09) and provided a fully characterized impact. Correlation analysis between mechanical inputs and degree of superficial cracking made visible through India ink application provided significant positive correlations for stress and stress rate with degree of surface cracking ($R^2 = 0.7398$ and $R^2 = 0.5262$, respectively). Ranges of impact parameters were 7 to 21 MPa, 6 to 40 GPa/s, 0.16 to 0.38, 87 to 236 s⁻¹, and 0.3 to 1.1 MJ/m³ for stress, stress rate, strain, strain rate, and strain energy density, respectively. Thresholds for damage for all inputs were determined at 13 MPa, 15 GPa/s, 0.23, 160 s⁻¹, and 0.59 MJ/m³ for this system. *Conclusions*: This study provided the mechanical basis for use of a portable, sterilizable, and maneuverable impacting device. Use of this device enables controlled impact loads *in vitro* or *in vivo* to connect mechanistic studies with long-term monitoring of disease progression.

Keywords

articular cartilage, in vitro, posttraumatic arthritis

Introduction

Osteoarthritis (OA) is the leading cause of chronic disability in the United States, and a significant fraction of OA cases are attributable to previous joint trauma (i.e., posttraumatic osteoarthritis [PTOA]).¹⁻³ No therapies currently exist that prevent or slow progression of PTOA, and mounting evidence suggests that interventions must occur in the acute time frame after injury to effectively modify the course of the disease.⁴ Therefore, research models that allow interrogation of pathologic mechanisms immediately following impact injury are critical to the development of preventative therapies. Both in vitro and in vivo impact models have been applied to native articular cartilage. Common methods to deliver impact injury include drop towers,⁵⁻⁷ pendulums,⁸ and more recently spring-loaded impactors.⁹⁻¹² Unlike stress or strain controlled impact systems,¹³⁻¹⁵ these impacting methods deliver a quick, single impact where energy input is correlated to the initial height of a weight or pendulum or to the stretch of a spring. With such systems, it is difficult to

fully characterize an insult mechanically as both stress and strain data are difficult to capture with such quick insults that occur during time scales on the order of milliseconds.

In previous work, an *in vivo* impactor⁹ consisting of a spring-loaded missile attached to a load cell was used to measure impact force, while pressure-sensitive film provided an estimate of contact area and impact stress. This device has major advantages for use *in vivo* including

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Figure 1. The spring-loaded impactor used in this study was modified from previous studies^{9,10} to include a displacement sensor (LVDT). This sensor was added in parallel with the impactor tip.

compact design (allowing sterilization and intraarticular positioning of the impactor tip) and rapid impact delivery. Limitations of the design include a lack of information on the deformations that occur during impact, as well as inertial effects of having a mounted load cell accelerate prior to impact that may confound real-time measurements of load during impact. In the present study, we have modified the impactor to more fully characterize the insult. Addition of a displacement sensor provided real-time displacement data. In addition to providing impact strain data, this allowed the calculation of the strain energy density, a possible metric for predicting surface damage.¹⁶

The goals of this study were 3-fold: (1) present a fully characterized insult that is capable of correlating stress, stress rate, strain, strain rate, and strain energy density to severity of damage (i.e., surface cracking); (2) provide a method to reduce inertial artifacts that occur when a load cell is mounted to a moving section of an impactor; and (3) determine which mechanical inputs correlate most strongly with measured surface damage.

Materials and Methods

Impactor Modification

A spring-loaded impactor presented by Alexander *et al.*⁹ was modified in this study to simultaneously provide stress and strain data (**Fig. 1**). In short, a 12-mm diameter

impacting head was mounted in line with the axis of a spring that was compressed to produce energy-controlled impacts. The impact force was measured at 50 kHz by a load cell (PCBPiezotronics, Depew, NY) attached to the impactor tip. A linear variable differential transducer (LVDT; RDP Electronics, Pottstown PA) was mounted in parallel with the impactor and attached to the impactor tip to measure the depth of penetration of the impactor tip into a sample in real time.

Sample Preparation and Analysis

In these validation studies, neonatal bovine cartilage was impacted and analyzed. Femoral condyle cartilage was explanted from 1- to 3-day-old bovids within 48 hours of sacrifice and sectioned into 6 mm diameter by 3 mm thick cylinders with the articular surface intact. Samples contained the majority of the deep zone tissue but no subchondral bone. Twelve samples were impacted under 6 different stretches of the internal spring, and voltages from the load cell and LVDT were recorded simultaneously with a custom LabVIEW program (NI, Austin TX). In the present study, a plane-ended impactor tip was used resulting in an impact configuration consistent with unconfined compression.

After impact, India ink was applied to the articular surfaces to detect superficial cracking and fibrillation. Consistently for all samples, several drops from a transfer pipette were applied to the surface and allowed to sit for no more than 3 minutes, after this time the samples were rinsed in phosphate-buffered saline (PBS) and wiped with a PBSsoaked Kimwipe to remove unbound dye. To quantify the amount of surface damage, photographs of the stained surfaces were imported into ImageJ (NIH, Bethesda, MD) and converted to binary images. The area percent of surface staining was recorded as a measure of surface damage.

Signal Analysis and Processing

Throughout all 12 impacts, the recorded voltage from the load cell resembled a 2-peaked impulse while the LVDT voltage resembled a 1-peak impulse. This 2-peak impulse, considered an artifact of accelerating the load cell along with the impacting head prior to impact, can systematically confound further mechanical analyses such as calculations of peak stress and stress rate. The first voltage peak is a result of the spring hitting the load cell and accelerating it downward. This portion of the impact is not directly applied to the cartilage surface. Here, we present a method of data analysis to remove the inertial artifact.

Assuming that the load cell signal is composed of 2 impulses, one peak when the load cell accelerated downward and one when the cartilage was impacted, the raw stress data were fitted to the sum of 2 Gaussian impulses given by the equation:



Figure 2. (**A**) The raw (unmodified) stress signal resembled a 2-peaked impulse for all impacts and fit well to the impact model (Equation 1). (**B**) The corrected stress signal fit well to a single impulse model (Equations 2 and 3). The constants *a* and *c* are shown for clarity. (**C**) Stress rate data of both the data and explicit derivative of the model (Equations 4 and 5). (**D**) Strain data fit well to a single impulse model (Equation 6). (**E**) Strain rate obtained similarly to stress rate. (**F**) Integration of the stress–strain curve provides a value for strain energy density.

$$\sigma(t) = a_1 \left(\exp\left(\frac{-(b_1 - t)^2}{2c_1^2}\right) \right) + a_2 \left(\exp\left(\frac{-(b_2 - t)^2}{2c_2^2}\right) \right) \quad (1)$$

where σ is the stress signal from the load cell, *a*, *b*, and *c* are constant fit parameters, and *t* is time. The constants *a*, *b*, and *c* correspond to the amplitudes of the impacts, the times when the impact peaks occur, and the durations (i.e., the standard deviation) of the impacts, respectively. The subscripts 1 and 2 correspond to the first and second impulses, respectively. The model constants were obtained by minimizing root mean square error (RMSE) between the model and data using a custom-developed Excel program (available on request). By subtracting the first impulse from the raw data, we obtained a corrected raw data set and corrected model curve:

$$\sigma_{\text{corrected}}(t) = \sigma_{\text{raw}}(t) - a_1 \left(\exp\left(\frac{-(b_1 - t)^2}{2c_1^2}\right) \right)$$
(2)

$$\sigma_{\text{model}}\left(t\right) = a_2\left(\exp\left(\frac{-\left(b_2 - t\right)^2}{2c_2^2}\right)\right)$$
(3)

The corrected data and model fit (Equations 2 and 3) are shown in **Figure 2B** where the parameters a and c from Equation (3) are highlighted for clarity. Obtaining a corrected data set and model fit allowed for calculation of stress rate both numerically and from differentiation of the

model (**Fig. 2C**). The equations for both the numerical and explicit stress rates are given by

$$\frac{d\sigma}{dt} = \frac{\sigma_t - \sigma_{t-5}}{5\Delta t} \tag{4}$$

$$\dot{\sigma}(t) = \frac{d\sigma(t)}{dt} = a_1 \frac{(b_1 - t)}{c_1^2} \left(\exp\left(\frac{-(b_1 - t)^2}{2c_1^2}\right) \right)$$
(5)

where a numerical derivative is taken over 5 data points to minimize noise and an explicit derivative of the model is taken with respect to time. Similar precautions were not necessary for strain and strain rate taken from the LVDT due to the 1-peak nature of the raw data. The strain model was similar to the stress model in Equation (3) and is given by

$$\varepsilon_{\text{model}}(t) = a_3 \left(\exp\left(\frac{-(b_3 - t)^2}{2c_3^2}\right) \right)$$
(6)

where ε is strain and a_3 , b_3 , and c_3 are distinct constants. Strain rates were then calculated in a similar way to stress rates.

Impact Strain Energy Density

With both stress and strain data available for each impact, the data were parametrically presented on a stress-strain



Figure 3. Typical model traces for (**A**) stress, (**B**) strain, (**C**) stress rate, (**D**) strain rate, (**E**) stress–strain, and (**F**) strain energy density. The letters a, b, and c denote data corresponding to impacts conducted at 3 different deflections of the internal spring.

curve. With this relationship, the strain energy density applied during an impact may be calculated from the following equation:

$$e = \int_{0}^{\varepsilon_{\max}} \sigma d\varepsilon \cong \sum_{\epsilon=0}^{\varepsilon_{\max}} \sigma_t * (\varepsilon_t - \varepsilon_{t-1})$$
(7)

where e is the strain energy density (energy applied per unit volume), ϵ is strain, and $\epsilon_{_{max}}$ is the maximum strain during an impact. For the impacts in this study, a right rectangular approximation of the integral was used to determine the amount of energy applied during an impact. The strain energy density is shaded in the stress strain curve for clarity (Fig. 2F). Due to the high frequency of data collection and using different sensors for stress and strain, the data were inconsistently out of phase, although the strain rates would imply an elastic response due to deformations greatly exceeding the gel diffusion rate in cases where there is minimal energy dissipation through viscous losses or damage initiation^{13,17} (Peclet number $\sim 10^6$). Due to this drawback in data acquisition, an elastic response was modeled where the peak stress and strain were assumed to occur at the same time.

Results

A representative fit of Equation (1) is presented in **Figure 2A**, where raw data are dotted, the full fit is the solid line, and the 2 Gaussian impulses are shaded in the background. Representative data and fits for stress, stress rate, strain,

strain rate, and stress-strain are shown in Figure 2B-F, where strain energy density is the shaded area in Figure 2F. Impacts with varying spring deflections provided a range of stress, stress rate, strain, strain rate, and impact energy values (Fig. 3). Peak impact stresses ranged from 7 to 22 MPa, with peak stress rates ranging from 6 to 40 GPa/s. The raw data fit well to the impulse models with average (\pm standard deviation, n = 12) coefficients of variation of the RMSE equal to 0.087 ± 0.028 and 0.103 ± 0.041 for the fits of Equations (1) and (3), respectively. Similarly, peak strains ranged from 0.16 to 0.38, and peak strain rates ranged from 87 to 240 s^{-1} . As with the stress data, all of the strain data fit well to the impulse models with average coefficient of variation of the RMSE equal to 0.084 ± 0.032 . Relating stress and strain through the energy density relationship (Equation 7), peak strain energy density varied from 0.302 to 1.08 MJ/ m'. Both peak stress and peak strain had a significant positive correlation with strain energy density (P < 0.05, data not shown). Furthermore, in 8 independent impacts at the same spring deflection, the coefficients of variation for stress, stress rate, strain, strain rate, and strain energy density were 0.066, 0.11, 0.088, 0.045, and 0.062, respectively (Fig. 4, n = 8).

Image analysis of the articular surfaces revealed significant cracking (arbitrarily determined at >1% staining by area) in samples with higher energy impacts. No significant surface damage was observed for peak stresses, stress rates, strains, strain rates, and strain energy densities below 13 MPa, 15 GPa/s, 0.23, and 160 s⁻¹, and 0.59 MJ/m³, respectively (**Fig. 5**).

Figure 4. The repeatability of the impactor was shown through independent impacts at one spring deflection (n = 8, bars denote mean).

Stress and stress rate correlated significantly with surface damage (**Fig. 5**, P < 0.05), while strain, strain rate, and strain energy density did not. These fits predicted damage thresholds of 14.4 MPa and 13.9 GPa/s for stress and stress rate, respectively, for this impacting system and tissue.

Discussion

This study describes the characterization of mechanical consequences of impact injuries to articular cartilage delivered with a hand-held instrument. This instrument, which has been previously used to deliver impact injuries *in vivo*,⁹ delivers impacts with peak stresses up to 20 MPa on a time scale of 2 ms, producing stress rates up to 40 GPa/s and macro-scale strains and strain rates of 0.4 and 200 s⁻¹, respectively. Use of this instrument enabled the identification of thresholds of mechanical inputs above which visible fissures were induced in the articular surfaces of cartilage explants. The mechanical factors that most strongly correlated with the development of tissue cracks were stress and stress rate.

The characterization of this model enables more mechanistic studies of PTOA *in vivo*. The ability to easily move and sterilize the impactor allows it to be used within the operating room. Furthermore, the ability to perform the impacts at arbitrary angles, as opposed to the vertical angles necessary for drop towers and horizontal angles necessary for pendulum systems, makes this system easier to use for *in vivo* studies where animal positioning may become a limiting factor. In addition to these innovations, the impacts have been shown to be fully characterized and controllable. This system enables controlled impacts *in vitro* and *in vivo* that may enable direct comparisons of mechanistic studies with long-term monitoring of disease progression.

The fully characterized mechanical insult resulted from the addition of a displacement sensor that provided the basis for measuring strain, strain rate, and also strain energy density. By correlating all of the mechanical aspects of the impact with observed surface damage, we unexpectedly found that only stress and stress rate significantly correlated with degree of surface damage. Although the correlations provided threshold values for only stress and stress rate, threshold values for strain, strain rate, and strain energy density were evident in the surface cracking plots (Fig. 5). Due to inherent variability between samples it is likely that strain, strain rate, and strain energy density are not sensitive metrics for predicting degree of surface damage. In addition to intersample variation, the finding that stress and stress rate are more predictive of damage than strain, strain rate, or strain energy density may be because they are more directly related to internal fluid pressure, which is known to be an important factor in the failure of poroelastic/biphasic materials through hydraulic fracturing.¹⁸ Consistent with other studies,¹³ this study suggests that stress and stress rate are the most sensitive metrics for predicting cartilage surface damage.

In the context of other studies, this device provides a set of impact parameters that fall in range of those previously reported. Other spring-loaded systems estimated peak stresses of 17 to 80 MPa, but did not provide data on stress rates or tissue strains.¹² In a drop tower system peak stresses of 60 MPa¹⁹ were observed while a strain-controlled system provided peak stresses of 14 MPa.¹³ Between those studies, peak strain rates ranged from 1000 s⁻¹ to 0.07 s⁻¹, respectively. Similarly, stress rates from other studies have varied from 35 MPa/s to 2.5 GPa/s for a strain controlled system and a pendulum system, respectively,^{8,20} but stress rate was not reported for the drop tower study. Previous in vitro studies using this system reported correlations between peak stress, cell death, proteoglycan release, and surface fissuring in a range similar to the one reported here (~17 MPa peak stress).¹⁰ It should be noted, however, that contact geometry and boundary conditions can also affect comparisons, where the results in this study utilized explants with no subchondral bone and are impacted in unconfined compression may differ from other configurations.

The stress and stress rate values determined in this study were obtained from fitting the data to a modified Gaussian impulse model. This model (Equations 1-5) was used to deconvolve inertial and impact signals from the mounted load cell. *In vivo* impact systems that operate using a mounted load cell that accelerates during impact may also have an inertial artifact present during the impulse force trace. The method used here corrects for this inertial artifact allowing calculation of stresses and stress rates that are not confounded by inertial effects. Simply put, the inertial artifact would overestimate the strain energy density and may also result in higher stresses and stress rates in cases where





Figure 5. Twelve samples were impacted at 6 different deflections of the internal spring. (**A**) Representative photographs and binary images of the 6 mm diameter surface for low (below damage threshold) and high impact (above damage threshold) samples after application of India ink. (**B-F**) Correlations between surface cracking and mechanical aspects of impact were determined from closed circle data points, which included all data points with >1% cracking and the highest point that did not reveal cracking. Only stress (**B**) and stress rate (**D**) provided significant correlations with surface cracking (P < 0.05).

the inertial peak exceeds or occurs faster than the tissue impact peak, respectively.

This study suggests that stress and stress rate may be sufficient to predict severity of impact after full characterization of an impact system. Future *in vivo* studies utilizing this system and analysis framework may reduce the current knowledge gap surrounding the mechanical and biological aspects of PTOA and provide information necessary to develop preventative strategies following cartilage injury.

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Declaration of Conflicting Interests

The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Ethical Approval

No live animals were used in this study. It was exempt from IACUC approval.

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