

Research Paper

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Intracranial Pressure Response to Non-Penetrating Ballistic Impact: An Experimental Study Using a Pig Physical Head Model and Live Pigs

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Abstract

This study was conducted to characterize the intracranial pressure response to non-penetrating ballistic impact using a "scalp-skull-brain" pig physical head model and live pigs. Forty-eight ballistic tests targeting the physical head model and anesthetized pigs protected by aramid plates were conducted with standard 9 mm bullets at low (279-297 m/s), moderate (350-372 m/s), and high (409-436 m/s) velocities. Intracranial pressure responses were recorded with pressure sensors embedded in similar brain locations in the physical head model and the anesthetized pigs. Three parameters of intracranial pressure were determined from the measured data: intracranial maximum pressure (P_{max}), intracranial maximum pressure impulse (PI_{max}), and the duration of the first positive phase (PPD). The intracranial pressure waves exhibited blast-like characteristics for both the physical model and I live pigs. Of all three parameters, P_{max} is most sensitive to impact velocity, with means of 126 kPa (219 kPa), 178 kPa (474 kPa), and 241 kPa (751 kPa) for the physical model (live pigs) for low, moderate, and high impact velocities, respectively. The mean PPD becomes increasingly short as the impact velocity increases, whereas Plmax shows the opposite trend. Although the pressure parameters of the physical model were much lower than those of the live pigs, good correlations between the physical model and the live pigs for the three pressure parameters, especially P_{max} , were found using linear regression. This investigation suggests that P_{max} is a preferred parameter for predicting the severity of the brain injury resulting from behind armor blunt trauma (BABT).

Key words: Intracranial pressure response; Non-penetrating ballistic impact; Pig physical head model; Live pigs

Introduction

Although ballistic helmets can prevent the penetration of ballistic projectiles, bullets, or shrapnel into the head, damage to the head caused by non-penetrating ballistic impact resulting from rapid deformation of the helmet is still difficult to prevent completely [1,2]. Non-penetrating injuries to the head behind a protective covering, which are included in behind armor blunt trauma (BABT), have not been researched as extensively as thoracic BABT [3]. Nonetheless, the BABT head injury pattern has been observed on human cadaver heads and dry skulls protected by polyethylene or aluminum plates when impacted by a non-penetrating 9 mm round [4,5]. As revealed by Sarron's reports, non-penetrating ballistic impact to the head while wearing protection resulted in skin lacerations, extensive skull fractures and brain damage, which were associated with increased pressure in the brain. This evidence suggested that intracranial pressure waves are the likely mechanism behind such injuries. However, the intracranial pressure response of brain tissue to non-penetrating ballistic impact resulting from BABT has been poorly documented to date.

The aim of this study was to characterize intrapressure cranial on brain tissue during non-penetrating ballistic impact to provide a better understanding of the biomechanics of BABT of the head. A physical model of the pig head and live pigs were used as objects of ballistic tests to acquire pressure data. Three characteristic parameters of the pressure wave were considered: intracranial maximum pressure, intracranial maximum pressure impulse, and the duration of the first positive phase. The sensitivity of pressure parameters to impact velocity was examined. Moreover, the intracranial pressure in the physical head model was compared with that in the live pigs.

Materials and Methods

Preparation of the pig physical head model and experimental animals

A "scalp-skull-brain" pig physical head model was developed to measure the intracranial pressure response to non-penetrating ballistic impact (Fig. 1). The pig physical head was molded from the approximate geometry of a white pig weighing 45.36 kg, which was provided by the Experimental Animal Center, Daping Hospital, Third Military Medical University (Chongqing, China). The scalp was made of polyurethane elastomer, while the brain was produced with silicone gel. Thermosetting resin mixed with calcium phosphate and fiberglass was used as the skull material.

Twenty-four white pigs weighing 43 ± 3 kg were divided randomly into three groups (n=8) for intracranial pressure tests for ballistic impacts at low, moderate, and high velocities. Animals were anaesthetized by transvenous injection of pentobarbital sodium (40 mg/kg) at the beginning of the experiment. All procedures were performed in accordance with the Guide for the Care and Use of Laboratory Animals as adopted by the Experimental Animal Center, Daping Hospital, Third Military Medical University.



Fig. 1. The pig physical head model.

Ballistic impact

Aramid plates ($200 \times 150 \times 9$ mm) were used to represent helmet protection and induce BABT in a pig physical head model and living pigs. To simulate the realistic space between the head and a ballistic helmet, a 12 mm gap between the protective plate and the subject was created by placing two pieces of foam padding, which is used specifically for helmets, on the nasion and the occiput of each subject. The nominal target location on each test specimen was the midpoint on the line drawn laterally between the eyes. The subject with ballistic protection was held in a lateral position, ensuring that the firing axis was perpendicular to the protective plate (Fig. 2). The ammunitions used for ballistic impact were 9 mm bullets at low, moderate, and high velocities of approximately 280 m/s, 360 m/s, and 420 m/s, respectively. These velocities were selected from preliminary experiments to provide varying injury risks with no chance of penetrating the protective plate that was

aligned with the nominal impact location on the subject. The ballistic impacts were conducted eight times at each velocity (low, moderate, and high) on both the physical head model and the live pigs (each live animal was impacted only once). Each plate was used only once in all tests.

Protective plate Firing axis 9mm Bullet

Fig. 2. Schematic representation of the ballistic impact scenario for both the pig physical head model and live pigs.

Recording of intracranial pressure

А piezoresistive pressure sensor (EZM-093-500(sg), Saiying Electronics, Anhui, China) was installed inside the brain of each test specimen both in the physical head model and in the anesthetized pigs (Fig. 3). The sensor was inserted into the brain parenchyma through a occipital approach and positioned approximately 12 mm under the parietal bone, facing the point of impact. The location of the sensor was controlled by a scheduled operation procedure performed by one person to ensure as much uniformity as possible among individual subjects. Signals were acquired using a TST6150 numerical recording system (Test Electronics, Chengdu, China) and analyzed with DAP6.01. The data were used to calculate the maximum value of the first positive phase of the post-impact curve of intracranial pressure versus time (P_{max}) , the duration of the first positive phase (PPD), and the integral of the intracranial pressure curve over the duration of PPD (PI_{max}).

Data analysis

Statistical analyses of intracranial pressure parameters were performed using SPSS 17.0 statistical software. One-way analysis of variance (ANOVA) was conducted on P_{max} , PI_{max} , and PPD for compari-

Fig. 3. Illustration of the location of the pressure sensor in the brain of a test specimen.

Results

Intracranial pressure of the physical head model and the live pigs

Intracranial pressure time histories were recorded for each ballistic test. As shown in Fig. 4 AB, the typical waveforms captured by the pressure sensors in the brain of the pig physical head model and live pigs exhibited blast-like features characterized by a steep shock front followed by a near-exponential decay. The waveforms of intracranial pressure corresponding to the different ballistic impact velocities display similar agreement.

The calculated P_{max} , PI_{max} , and PPD values for each test are included in Table 1. For all three pressure parameters, the values for the physical head model were significantly lower than those for the live pigs. The P_{max} means were 126 kPa, 178 kPa, and 241 kPa for the physical head model and 219 kPa (p<0.01), 474 kPa (p<0.01), and 751 kPa (p<0.01) for the live pigs under low, moderate, and high impact velocities, respectively. The PPD was also significantly different between the pig physical head model and the live pigs, with means of 0.039 ms, 0.032 ms, and 0.028 ms for the physical head model and 0.311 ms (p<0.01),



son between different impact velocities. A probability value of <0.05 was considered significant. In addition,

linear regressions were performed to determine not

only the sensitivity of intracranial pressure parame-

ters to ballistic impact velocity but also the correlation

of all three pressure parameters between the physical

head model and the live pigs.

0.238 ms (*p*<0.01), and 0.181 ms (*p*<0.01) for the live pigs under low, moderate, and high impact velocities, respectively. As the maximum value of the integral of the intracranial pressure curve within PPD, PI_{max} showed magnified differences between the pig physical head model and the live pigs relative to PPD, with means of 1.83 kPa ms, 3.46 kPa ms, and 4.54 kPa ms for the physical head model and 64.50 kPa ms (p<0.01), 88.11 kPa ms (p<0.01), and 132.78 kPa ms (p<0.01) for the live pigs under low, moderate, and high impact velocities, respectively.

Table I. Intracranial pressure records

Target	Impact velocity	Pmax (kPa)	PImax (kPa ms)	PPD (ms)
	(m/s)			
physical model/ low velocity				
1	279	112	1.65	0.025
2	287	107	1.70	0.044
3	292	119	1.53	0.026
4	297	138	2.06	0.034
5	296	149	2.31	0.030
6	295	125	1.76	0.058
7	289	128	2.03	0.045
8	297	130	1.62	0.046
physical model/ moderate veloc- ity				
9	366	188	3.84	0.036
10	356	172	3.87	0.030
10	355	171	5.04	0.029
12	359	185	4 59	0.047
13	350	171	2.19	0.027
14	361	167	2.36	0.024
15	360	165	3.21	0.022
16	365	203	2.59	0.022
physical model/ high velocity				
17	409	215	3.28	0.023
18	411	210	4.99	0.020
19	429	279	3.97	0.054
20	417	218	7.16	0.048
21	420	238	3.38	0.017
22	419	250	5.15	0.024
23	426	253	4.39	0.019
24	422	268	3.96	0.015
live pigs/ low velocity				
25	282	178	51.27	0.257
26	295	241	47.79	0.202
27	289	190	82.48	0.665
28	280	182	82.15	0.143

30	293	289	48.27	0.184
31	296	269	74.29	0.506
32	288	193	79.64	0.283
live pigs / moderate veloc- ity				
33	371	583	134.49	0.446
34	366	424	70.45	0.093
35	370	439	70.03	0.164
36	372	566	59.50	0.143
37	368	532	120.88	0.269
38	360	404	84.68	0.158
39	365	416	72.17	0.437
40	363	428	92.68	0.192
live pigs / high velocity				
41	422	705	153.29	0.218

212

50.08

291

431

434

419

429

430

436

420

29

> 41 42

43

44

45

46

47

48

Sensitivity of pressure parameters to impact velocity

892

832

653

701

740

837

647

149.72

148.52

153.35

138.03

148.19

100.53

70.63

ANOVAs were used to compare pressure parameters (Pmax, PImax, PPD) at different ballistic impact velocities. Data from the physical head model and live pigs were examined. The P_{max}, PI_{max}, and PPD values obtained from the physical head model and live pigs are plotted against impact velocity in Fig. 5A,B,C. Of the three parameters, P_{max} was the most sensitive to impact velocity, corresponding to the ANOVA finding that differences in P_{max} between different impact velocities are significant (p < 0.01) for both the physical head model and live pigs. PI_{max}, which exhibited significant differences between different ballistic impact velocities, increased with the impact velocity. However, the PPD values under different impact velocities did not show statistically significant differences (p>0.05) for either the physical head model or the live pigs, although there was a non-significant decrease in PPD with increasing impact velocity. The intracranial pressure response of the live pigs is more sensitive to impact velocity compared with that of physical head.

Correlations

The correlation parameters between the physical head model and live pigs for intracranial pressure

0.251

0.203

0.177

0.092

0.247

0.176

0.132

0.203

parameters are provided in Table 2. For each correlation, data from the physical head model and live pigs under low, moderate, and high impact velocities were used. Fig. 6A,B,C show the correlations of P_{max} , PI_{max} , and PPD between the physical head model and live pigs. Of the three pressure parameters, P_{max} of the physical head model correlated best with that of live pigs (R^2 =0.9785, p<0.001). **Table 2.** Correlations of intracranial pressure between thephysical head model and live pigs (N=24)

	R ²	р	
P _{max}	0.9785	< 0.001	
PI _{max}	0.8769	< 0.001	
PPD	0.8942	<0.001	



Fig. 4. Intracranial pressures in the brain of the pig physical head model(A) and a live pig(B) with a protective aramid plate following the impact of 9 mm bullets at low, moderate, and high velocities.



Fig. 5. Intracranial maximum pressure $(P_{max})(A)$ and Intracranial maximum pressure impulse $(PI_{max})(B)$ of the pig physical head model and the live pigs for different impact velocities. Duration of the first positive phase (PPD) of the intracranial pressure wave in the pig physical head model and the live pigs for different impact velocities(C).



Fig. 6. Correlation of the intracranial maximum pressure $(P_{max})(A)$ and intracranial maximum pressure impulse $(PI_{max})(B)$ between the pig physical head model and live pigs. Correlation of the duration of the first positive phase (PPD) of the intracranial pressure wave between the pig physical head model and live pigs(C).

Discussion

Non-penetrating injuries to the head resulting from BABT have been studied in human cadavers, dry skulls filled with silicone [4,5], and finite element (FE) models of the human head [6]. Although these types of models have accurate anatomy or similar anthropometry to humans, their lack of living physiology remains a main limitation for further investigation of the post-injury physiological response to BABT. Animals possessing living physiology are the only viable subjects in which to study the medical outcomes of non-penetrating injuries to the head, including not only anatomical features but also physiopathological changes, and can therefore help correlate biomechanical response data with the resulting BABT injuries to the head. In our study, we used pigs as ballistic subjects and focused on the brain's intracranial pressure response to non-penetrating impacts to the protected head. In addition, a pig physical head model was also used because of its reusability, which may contribute to providing comparatively repeatable response data. As previously reported, BABT injury to the head may be characterized physically by skin lacerations, skull fractures, and brain damage. Many authors indicate that the skin and scalp absorb 13% to 20% of incident energy. A study of transcranial transmission of blast waves revealed that blast waves pass through the thicker cranium of pigs while maintaining about two-thirds of their original magnitude [7]. Thus, the structural configuration of the "scalp-skull-brain" for the pig physical head model developed in this study is suitable for investigating the biomechanical response of BABT to the head.

High intracranial pressure was observed in the brain as a result of exposure to non-penetrating ballistic impact. However, the pressure waves propagating through the tissue have some of the characteristics of blast waves. Because of these pressure waves, the peak pressure, duration, and impulse can be regarded as contributors to injury [8]. Thus, three parameters, P_{max}, PI_{max}, and PPD, were extracted from the recorded pressure curve to describe the characteristics of the intracranial pressure response. P_{max} is the maximal instantaneous pressure recorded in the brain, corresponding to local compression of the surrounding tissue under skull deflection. PImax corresponds to the first positive area under the intracranial pressure curve, and refers to the amount of energy transferred to the head. It depends not only on P_{max} but also on PPD, which is defined as the duration of the first positive phase. In a previous study on the dynamic effects of projectiles on cadaveric skulls protected by a helmeted plate, the risk of skull fractures correlated well with brain parenchymal pressures [5], indicating that intracranial pressure is an appropriate predictor for studying the effects of cephalic impacts behind helmets. Injury criteria based on intracranial pressure have not yet been determined for BABT injuries to the head. However, conclusions from the related research areas of blast-induced traumatic brain injury (bTBI) and thoracic BABT are also instructive. A study focusing on the biomechanical assessment of brain response to blast pressure waves suggested that the maximum intracranial pressure can be compared with the threshold injury pressure and used to assess the severity of brain injury and the occurrence of cerebral contusion [9]. The intrathoracic pressure impulse was also found to be a useful predictor for pulmonary contusion volume after ballistic blunt impact thoracic trauma [10]. Further investigations are needed to clarify the function of intracranial pressure as an injury criterion for BABT injuries to the head.

To obtain a large range of intracranial pressure responses, both the physical head model and live pigs were exposed to ballistic impacts of standard 9 mm bullets at low (279-297 m/s), moderate (350-372 m/s), and high (409-436 m/s) velocities. Non-penetrating ballistic impacts produced lower intracranial pressures (mean Pmax=126 kPa, 178 kPa, and 241 kPa for low, moderate, and high impact velocities, respectively) in the surrogate and higher intracranial pressures in the living specimens (mean P_{max}=219 kPa, 474 kPa, and 751 kPa for low, moderate, and high impact velocities, respectively). For comparison, Sarron's study [5] conducted on helmeted human cadaveric skulls reported intracranial pressures of 892±222 kPa after impact by non-penetrating 9 mm rounds at a velocity of 410.1±5.5 m/s. The higher intracranial pressures measured in this study may have resulted from the lack of scalp and anatomical differences in the human cadaveric skull subjects. Moreover, P_{max} for both the pig physical head model and live pigs was quite sensitive to impact velocity, indicating that P_{max} is a promising parameter for predicting brain injury severity resulting from BABT.

The PPD characteristics of the intracranial pressure response to non-penetrating ballistic impact have yet to be detailed. However, the data from several other BABT studies can be used as a point of reference. Bass et al. [11] confirmed that the impact duration is less than 300 μ s when a helmet-head system is impacted by a 9 mm bullet. This impact duration is comparable to the PPD values of intracranial pressure in the brains of live pigs (mean PPD=311 μ s, 238 μ s, and 181 μ s for low, moderate, and high impact velocities, respectively) in our study. The mean PPD values of intracranial pressure in the brain of the physical head model are 42 µs, 32 µs, and 28 µs for low, moderate, and high impact velocities, respectively, which are much lower than those for the live pigs. This order of magnitude for PPD has been observed in the pressure responses of internal organs to non-penetrating ballistic impacts conducted on a physical human surrogate torso model (HSTM) [12]. According to the description of the HSTM, the internal organs were made of silicone gel, which was similar to that used to make the brain of pig physical head model. Thereby, the similarity in the material composition is likely to be the main cause of the similarity of PPD values of pressure response in the HSTM and pig physical head model. Despite the fact that PPD decreased with increasing impact velocity, statistical analysis revealed that PPD was relatively insensitive to impact velocity for both the physical head model and live pigs. It is quite likely that the differences between the impact velocities are too small to produce statistically significant differences in PPD. Considering that the PPDs for the live pigs were much shorter than the overpressure durations of short-duration blasts (< 30 ms), which has a substantial effect on the peak overpressure tolerance, the exceptionally short PPD of the intracranial pressure waves will most likely be only a minor factor in BABT head injuries.

The difference in PI_{max} between the pig physical head model and the live pigs was increased by integral calculation of intracranial pressure vs. time during PPD. However, the PI_{max} for the live pigs was more sensitive to impact velocity than that for the pig physical head model.

Although considerable differences existed in the intracranial pressure parameters between the pig physical head model and the live pigs, especially in PI_{max} , linear regression has revealed correlations in the intracranial pressure parameters between the two subject types. Of the three pressure parameters, the P_{max} of the physical head model correlated best with that of live pigs. This finding suggests that it is feasible to predict the value of P_{max} in a live pig using its physical surrogate.

As we know, human surrogates have seen increasing use in obtaining mechanical responses to blunt ballistic impacts, which could improve the prediction of injury and the evaluation of protective equipment performance against projectiles [13]. However, discrepancies in the mechanical responses of surrogates compared with humans are inevitable owing to the simplification of human anatomic structure and differences between the material properties of biosimulants and living tissue [14]. Therefore, knowledge of the correlation between the mechanical responses for human surrogates and live pigs is critical for the better understanding and appropriate usage of surrogate response data. Unfortunately, such information remains unavailable because response data pertaining to humans in physiological conditions are understandably inaccessible. Animals, long used as live pigs for ballistic tests, are the only viable subjects for studying a living body's mechanical responses to ballistic impact and therefore contribute to interpreting the biomechanical correlation between physical surrogates and live pigs. In this study, we developed a physical model of the pig head to measure intracranial pressure responses to non-penetrating ballistic impacts and investigate the correlation between the physical surrogate and living specimens. We hope our work may help predict the biomechanical responses of living animals to blunt ballistic impact based on surrogate tests and also provide insight into controlling for differences between living humans and surrogates when developing response and injury targets.

Despite the good correlation of Pmax between the physical head model and the live pigs, the limitation of the physical model in lack of viscoelasticity for the brain material induced considerable differences of PPD and PImax between the physical head model and live pigs. Of course, an effort should be made to adjust the material properties of the physical model to correct its response and provide better correlation with the live pigs.

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Competing Interests

The authors have declared that no competing interest exists.

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