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19 **Integration of biplanar X-ray, 3-D animation, and particle simulation reveals details of**
20 **human ‘track ontogeny’**

21

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42 **Abstract**

43 The emergence of bipedalism had profound effects on human evolutionary history but the
44 evolution of locomotor patterns within the hominin clade remains poorly understood. Fossil
45 tracks record the anatomy and kinematics of extinct hominins, and they offer great potential to
46 reveal locomotor patterns at various times and places across the human fossil record. However,
47 there is no consensus on how to interpret anatomical or biomechanical patterns from tracks due
48 to limited knowledge of the complex foot-substrate interactions through which they are
49 produced. Here we implement engineering-based methods to understand human track formation
50 and potentially unlock invaluable information on hominin locomotion from fossil tracks. We first
51 developed biplanar X-ray and 3-D animation techniques that permit visualisation of subsurface
52 foot motion as tracks are produced, and that allow for direct comparisons of foot kinematics to
53 final track morphology. We then applied the discrete element method to accurately simulate the
54 process of human track formation, allowing for direct study of human track ontogeny. This
55 window lets us observe how specific anatomical and/or kinematic variables shape human track
56 morphology, and it offers a new avenue for robust hypothesis testing in order to infer patterns of
57 foot anatomy and motion from fossil hominin tracks.

58

59 **Keywords:** hominin footprints, trace fossils, locomotion, discrete element method

60

61 **Introduction**

62 Central to the study of human evolution are questions concerning the evolution of our
63 unique form of bipedal locomotion. While bipedalism has long been considered a defining trait
64 of the hominin clade (1), discoveries within the past half-century have made it apparent that

65 multiple forms of bipedalism likely existed among fossil hominins. Some of these forms were
66 probably quite similar to our own bipedal locomotion but others were almost certainly quite
67 different (2). To date, most evidence for the inferred locomotor patterns of fossil hominins has
68 come from comparative morphological studies of postcranial skeletal fossils. However, fossil
69 hominin tracks (i.e., footprints) have augmented, and have the potential to further augment, these
70 comparative osteological studies in important ways.

71 Tracks offer the only data on whole-foot anatomy, foot posture, and foot kinematics in
72 fossil hominins. Fossil hominin foot bones are most often found in isolation and even the most
73 exceptional, “nearly complete” hominin foot skeletons are missing important elements (e.g., OH
74 8 [*Homo habilis* (3)]; LB1 [*Homo floresiensis* (4)]; Foot 1 [*Homo naledi* (5)]; DIK-1-1f
75 [*Australopithecus afarensis* (6)]). Tracks are morphological features that result from the dynamic
76 interaction between the composite foot morphology (articulated foot skeleton and its soft tissues)
77 and a deformable substrate. Understanding, or reverse-engineering this interaction means tracks
78 can offer a picture of extinct hominin foot morphology complimentary to that offered by the
79 bones alone. At the same time, tracks record the three-dimensional kinematics of feet as they
80 navigated deformable substrates (7), allowing one to observe foot postures and motion patterns
81 that were actually used during bouts of terrestrial bipedalism. While the articular surfaces of
82 skeletal fossils might provide rough estimates of maximal joint mobility (but see (8)), tracks
83 result from specific poses and motion sequences that can help one to understand how hominin
84 feet were actually used to accomplish particular forms of bipedal locomotion.

85 In addition to tracks being able to augment analyses of skeletal fossils in critical ways,
86 fossil hominin track sites have been discovered at a high rate in recent years. The known record
87 of hominin track sites that predate modern humans has experienced notable growth (9–15). In

88 some cases, the known sample sizes of hominin tracks now exceed by more than an order of
89 magnitude the sample of hominin foot skeletal fossils from the same time periods (12). New
90 technologies are also being applied to digitally record hominin tracks in 3-D, thereby opening
91 doors for digital preservation, data sharing, and computational analyses (16,17).

92 Yet despite the great potential of these data and numerous recent advances in hominin
93 ichnology, there still exist major obstacles that limit access to the invaluable information
94 preserved by fossil hominin tracks. Perhaps the most important obstacle is our currently limited
95 understanding of the complex interactions between foot anatomy, kinematics, and substrate
96 through which a track is formed (18–20). Morse et al. (21) demonstrated, through a case study of
97 Holocene human tracks from Namibia, that track morphology can vary substantially as the same
98 individual walks through substrates of different consistencies. Yet the underlying reasons for that
99 variation remain unknown. Deciphering the mechanical nature of foot-substrate interactions is
100 essential for linking aspects of track morphology to anatomical or kinematic patterns (19) and
101 thereby for leveraging hominin tracks to better understand the evolution of human foot anatomy
102 and locomotion.

103 Falkingham and Gatesy (22) coined the term “track ontogeny” to describe the mechanical
104 process through which tracks are formed. This term emphasizes the fact that track morphology
105 develops through a dynamic sequence of continuous interactions between foot and substrate.
106 This developmental sequence is inherently difficult to study because track creation is usually
107 hidden from view – both human feet and natural substrates are opaque and so their interactions
108 cannot be observed directly. Building upon earlier biomechanical and robotic studies that used
109 X-rays to visualize subsurface motion (e.g., 23), Ellis and Gatesy (24) and Falkingham and
110 Gatesy (22) introduced biplanar X-ray approaches for studying 3-D foot-substrate interactions

111 that result in track formation. Those studies focused on track formation in guineafowl, but their
112 biplanar X-ray approach was more recently adapted and applied to study track formation in
113 humans (25).

114 Falkingham and Gatesy (22) were also the first to use particle simulation to understand
115 track ontogeny, by using the Discrete Element Method (DEM) to examine the mechanistic
116 origins of track morphology. The DEM simulates individual sediment particles as they interact
117 with each other and external geometry. These particle interactions are governed by physical
118 parameters including elasticity, compressibility, cohesion, and mass (26,27). By iteratively
119 simulating track formation processes, with consistent validation using experimental data,
120 Falkingham and Gatesy (22) and Falkingham et al. (28) were able to leverage their ontogenetic
121 perspective to develop robust inferences of trackmaker foot anatomy and foot kinematics from
122 fossil dinosaur tracks.

123 Here, we present the development and first application of similar methods that employ
124 biplanar X-ray, 3-D animation, and particle simulation to study track ontogeny in humans
125 walking through deformable muds. We build on existing methods in important ways, most
126 notably by animating and simulating high-resolution deformable 3-D models of human feet as
127 they interact with deformable substrates. We present a case study in which we demonstrate the
128 application of new methods, and potential directions for future research. These methods allow us
129 to open the black box of the foot-substrate interactions through which tracks are formed, and
130 they provide an avenue for robust inferences of foot anatomy and kinematic patterns to be
131 derived from fossil hominin tracks.

132
133

134 **Methods**

135 *Biplanar X-ray experiments*

136 Subjects

137 The methods presented here were developed and applied through experiments with four
138 healthy adult volunteer subjects, though as a proof of concept we present focused analyses from
139 only one individual. Subjects were recruited and provided informed consent to participate
140 through protocols approved by the Institutional Review Boards of Chatham University and
141 Brown University.

142

143 Biplanar X-ray setup and technique

144 The biplanar X-ray equipment, and its configuration within the W.M. Keck Foundation
145 XROMM Facility at Brown University closely followed that used by Hatala et al. (25). Details
146 on this configuration and recording settings are provided in Supplementary Text S1.

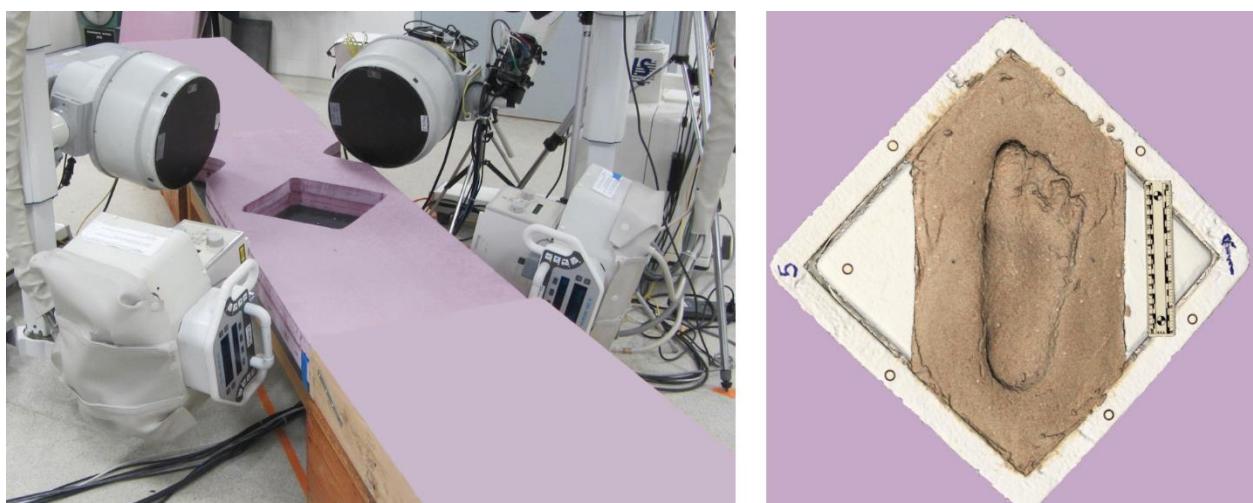
147

148 Trackway and substrates

149 A roughly 6-meter long (~60 cm wide, ~50 cm tall) elevated trackway was assembled,
150 following a setup that we have used previously to study human track formation via biplanar X-
151 ray (25). The biplanar X-ray apparatus was configured at roughly the center of this trackway,
152 with the two X-ray beams at an angle of approximately 90 degrees to each other. To improve
153 visibility of markers on the sole of the foot, the X-ray beams were pitched upwards 10 degrees
154 relative to the ground plane. X-ray emitters and image intensifiers were placed with a source-to-
155 image distance of 134 cm. X-ray videos captured anteromedial and anterolateral projections of
156 subjects feet.

157 The trackway was configured such that different substrates of interest could be placed
158 within the area of biplanar X-ray overlap. A modified stone slab table formed a rigid and stable
159 base within this central portion of the trackway. Three rigid, closed-cell foam panels (two 2-
160 inches thick, one 1-inch thick) were placed on top of the stone slab, and a diamond-shaped recess
161 was cut in the center of them, providing a space in which an interchangeable substrate container
162 could be securely placed (Fig. 1).

163



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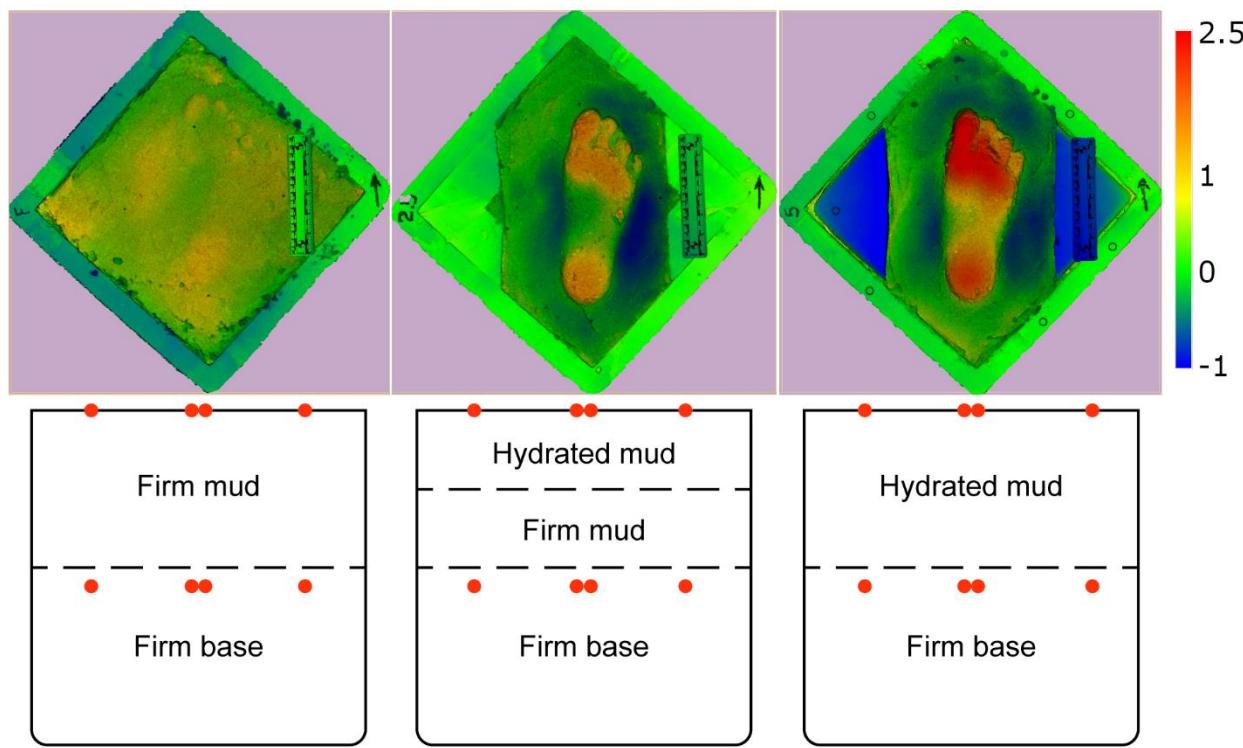
165 **Figure 1.** Edited photo showing trackway and biplanar X-ray configuration used in track
166 formation experiments. Portions of the trackway preceding and following the central, substrate-
167 bearing section were covered with various foams to make the entire trackway level and equally
168 deformable under each substrate condition. The central section includes a diamond-shaped recess
169 into which substrate containers were placed. The panel on the right shows an overhead view of a
170 3-D scan of the substrate container, with a track produced within it (in “hydrated 5” mud).

171

172 This configuration allowed for the study of foot motion on four substrates. In one setup, a
173 rigid foam core carbon fiber panel (79 x 30.5 x 2.7 cm) was placed over top of the recess, and 1-

174 inch closed-cell foam panels were placed along the remaining length of the trackway in order to
175 make it level. In the remaining three setups, a square foam container (30 cm by 30 cm opening,
176 14.5 cm deep, with 3 cm walls) was placed within the recess. Foam wedges were placed in the
177 medial and lateral corners of the substrate container, in order to reduce the volume of
178 “unnecessary” mud that X-rays would have to traverse but that would not interact with the foot
179 (thereby improving clarity of the X-ray videos). This left an area 22 cm wide, which held one of
180 three varieties of mud into which the foot would impress (Fig. 2). In these configurations, the
181 remainder of the trackway was topped with panels of rigid, closed-cell foam (for “firm” mud,
182 described below) or soft, deformable upholstery foams (approximately 2.5 cm thick for
183 “hydrated 2.5” mud, 5 cm thick for “hydrated 5” mud, described below) to mimic the
184 deformative natures of the substrates of interest and provide a level surface along the entire
185 trackway length.

186



187

188 **Figure 2.** Side-by-side comparisons of 3-D track models from the same subject in the three
189 varieties of mud (top row), alongside schematics showing the contents of substrate containers
190 (bottom row). Substrates included “firm” mud (left), “hydrated 2.5” mud (center), and “hydrated
191 5” mud (right). Track depth is reflected by color gradients according to scale at far right, which
192 is displayed in centimeters. Each substrate container included 6.5 cm of “firm base”, and an
193 overlying 5 cm that was filled according to the substrate conditions of that particular trial. At the
194 locations of orange dots, radiopaque marker beads were placed within and upon each substrate in
195 diamond-shaped patterns, to align the final track model within the same calibrated space as the
196 foot during 3-D animation.

197

198 Building upon previous biplanar X-ray studies of track formation (22,25,29), we
199 developed a new range of radiolucent, deformable, and cohesive muds that mimic the
200 mechanical behaviors and particle dimensions of naturally-occurring muds. These muds
201 consisted of 60 micron glass bubbles (Type K15, 3M Co., St. Paul. MN, USA), modeling clay,
202 water, and acrylic blast media (Type V, 0.42-0.56 mm diameter; Kramer Industries, Inc.,
203 Piscataway, NJ, USA). The first three ingredients were mixed in a 24:5:9 volumetric ratio
204 (following (29)) and this combination was then mixed with the acrylic blast media in roughly
205 equal volumetric proportions. In filling the substrate containers with mud, a substantial base
206 portion of substrate would not interact directly with subjects’ feet. In the bottom-most 6.5 cm of
207 substrate, we integrated EPS foam pellets (2-4 mm diameter; LACrafts) with the above
208 ingredients, to further enhance radiolucency while still maintaining relatively consistent material
209 properties throughout the substrate volume. Slightly beneath the surface of this firm base we
210 placed four radiopaque markers 3 mm in diameter, such that we could track those points and

211 identify and account for any potential disturbance to the entire substrate volume. The remaining
212 5 cm were then filled with one of three mud variants. In the “firm” mud condition, the substrate
213 container was filled to the rim with acrylic mud and tightly packed by tamping with a rubber
214 mallet. In the “hydrated 2.5” condition, 2.5 cm of “firm” mud was added atop the firm base.
215 Water was added to acrylic mud to make it more fluid and deformable and this filled the most
216 superficial 2.5 cm of the substrate container.. In the “hydrated 5” condition, the entire most
217 superficial 5 cm of the substrate container was filled with the hydrated acrylic mud. On the
218 surface of each of these substrates, we again placed four radiopaque beads 3 mm in diameter,
219 such that we could use those points to register the position of the final track during 3-D
220 animation (Fig. 2).

221

222 Experimental protocol

223 Subjects had an array of 85 radiopaque beads placed on the external surface of their right
224 foot, the motions of which could be tracked via biplanar X-ray. Some of these markers were
225 placed at anatomical locations of interest, but others filled in gaps to provide a roughly uniform
226 mesh of markers across the entire plantar surface of the foot. This array of bead markers expands
227 upon a 70 marker array used in earlier experiments (25) to achieve even more complete surface
228 coverage. Before marker beads were placed, a template was drawn on each subject’s foot using
229 semi-permanent marker. The foot was then 3-D scanned at 1.0 mm resolution using a handheld
230 structured light scanner (Creaform Go!SCAN 50, Creaform, Lévis, Québec, Canada; Fig. 3).

231 Following scanning of the foot with its marker template, 1.5-mm diameter radiopaque markers
232 (SureMark, Simi Valley, CA, USA) were placed and secured using medical adhesive
233 (SkinTacTM, Torbot, Cranston, RI, USA). After markers were placed, subjects moved to the

234 experimental trackway and walked across it several times until they were fully comfortable
235 moving within that environment.

236



237

238 **Figure 3.** High-resolution 3-D scan of a subject's foot with template for marker placements
239 drawn in semi-permanent marker. Views are plantar (center), lateral (left), medial (right), dorsal
240 (top). No markers were placed on the dorsum of the foot aside from those on the dorsal sides of
241 the toes.

242

243 Subjects traversed the experimental trackway for at least 13 trials each. In one trial, the
244 subjects simply stood with their right foot on the carbon fiber plate (with their left foot
245 immediately behind for support) while a single pair of X-ray images were taken of their
246 “statically loaded” marked foot. Each subject then walked across each of the four substrates
247 (carbon fiber and the three mud variants) for at least three trials at their self-selected comfortable
248 walking speed. If their foot strayed outside of the biplanar X-ray view, they were asked to repeat
249 that trial. For trials in which subjects walked through mud, the track they created was 3-D
250 scanned. For most trials the structured light scanner was used to scan the track at 1.0 mm
251 resolution. However, there were nine trials in which the scanning software was still processing
252 the model from the previous trial, and therefore we scanned tracks using photogrammetry
253 (Canon 5D Mark III camera, Canon, Melville, NY, USA; Agisoft Metashape Professional
254 v.1.6.4, Agisoft LLC, St. Petersburg, Russia). After track scanning, the substrate was
255 reconfigured to its initial state using a trowel, or swapped for a different substrate before the next
256 trial.

257

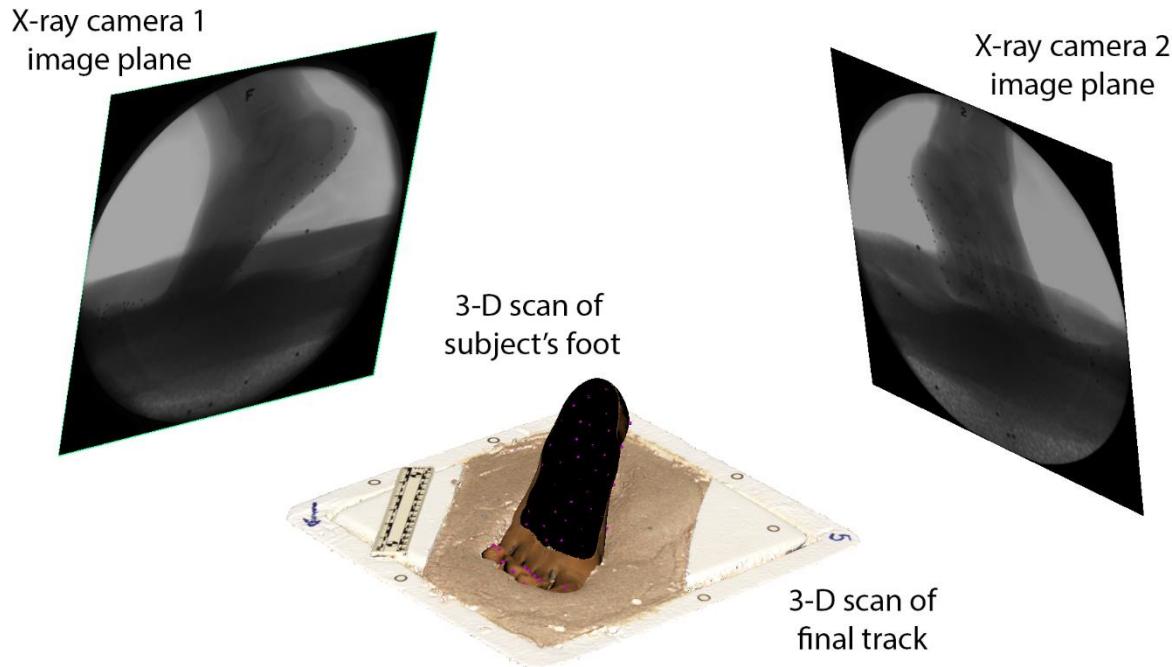
258 *Motion tracking and 3-D animations*

259 XMALab software (v.1.5.5) was used to compute the 3-D trajectories of radiopaque
260 marker beads that were placed on the foot, as it moved on and within the substrates of interest.
261 Following protocols that were established for X-Ray Reconstruction of Moving Morphology
262 (30,31), XMALab was used to remove distortion from video recordings, calibrate the 3-D
263 volume in which biplanar X-rays overlapped, and then track marker trajectories in 3-D. Since our
264 markers were placed on non-rigid human feet, and we sought to track soft tissue deformations
265 and motions, there was no informed basis for applying a filter to these data. Further, we used

266 XMALab's polynomial fitting procedure to improve sub-pixel accuracy (a procedure that has
267 been shown to reduce standard deviations of inter-marker distances on rigid bodies (31)), and
268 recorded at speeds of only 50 Hz, which should have the effect of minimizing potential "noise"
269 in 3-D marker trajectories. Additional details regarding marker tracking are provided in
270 Supplementary Text S2.

271 High-resolution scans of subjects' feet were processed and cleaned using Creaform
272 VXElements software (v. 7.0.1). Built-in mesh editing features were used to remove noisy
273 polygons (i.e., those discontinuous with the foot model) and to trim the foot model such that it
274 included, in general, only the area distal to the medial and lateral malleoli. These 3-D models
275 were exported in .obj format and then imported in Autodesk Maya 2020 for animation.

276 In the animation protocol, the high-resolution foot mesh was first imported to Autodesk
277 Maya 2020. For each individual trial, the 3-D coordinates of foot markers were imported into
278 Maya and animated as a collection of spheres each 1.5 mm in diameter using the "imp" function
279 of XROMM MayaTools (v. 2.2.3) (32). The positions of these spheres were linked to the
280 positions of the bead markers on the surface of the high-resolution foot model (Fig. 3;
281 Supplementary Text S3). The spheres were inter-connected such that their motions moved the
282 vertices of a low-resolution mesh, which in turn drove motions of the high-resolution mesh using
283 Maya's wrap deformer function (Supplementary Text S3, Supplementary Figure S1). Through
284 this series of connections and deformations, biplanar X-ray data were used to create trial- and
285 subject-specific animations of both aerial and sub-surface skin movements during track
286 formation (Fig. 4).



287

288 **Figure 4.** Snapshot of an animation of a single trial from biplanar X-ray experiments. The
 289 position of the mobile and deformable high-resolution 3-D foot scan is continuously guided by
 290 the tracked 3-D positions of external foot markers. Markers on the external surface of the foot
 291 appear as black dots in X-ray camera views, and are highlighted in purple for the sake of
 292 visibility on the animated foot model. The foot animation is integrated with a 3-D model of the
 293 final track that was produced in this trial, registered within the same calibrated 3-D space.
 294 Integration of feet and tracks within the same animation scene allows for direct visualization of
 295 the correspondence between track morphology and pedal kinematics.

296

297 Spheres (3.0 mm in diameter) were also animated to represent markers placed within and
 298 upon the substrate (Fig. 2). The final configuration of the four markers visible on the tracked
 299 surface were used to translate and rotate the scan- or photogrammetry-derived 3-D track model
 300 into registration. Such registration is critical for assessing the correspondence (or lack of
 301 correspondence) between pedal kinematics and track morphology. However, because only the

302 final track was captured, the integration of a dynamic foot with a static footprint (Fig. 4) is
303 insufficient to fully explain the origin and modification of specific features during a step. For
304 insights into the interplay between foot shape, foot motion, and substrate displacement, we
305 turned to simulation.

306

307 *Simulating track formation*

308 We used LIGGGHTS (www.cfdem.com; 27) to carry out discrete element simulations of
309 foot-substrate interactions. Our simulation process began with relatively simple foot motions and
310 iteratively increased motion complexity, in line with the animation process outlined above. All
311 simulations used the same initial particle set-up and parameters. A virtual tray 21 cm x 35 cm
312 and 8 cm deep was created in Autodesk Maya in the same world-space position as the original
313 substrate container. This completely encompassed the track-forming volume, though the virtual
314 tray lacked the diamond-shaped ends of the real substrate container for computational simplicity.
315 The virtual tray was filled with ~800,000 particles of 1 mm radius. While this particle size is
316 homogeneous and significantly larger than the experimental substrate, particle properties
317 (Young's modulus, Poisson ratio, cohesion, and friction) were adjusted such that the
318 macroscopic bulk behaviour was similar to our substrate.

319 The simplest simulation involved a vertical stamping of a rigid foot model (the scan of
320 the subject's foot in resting pose). Sinking depth of the rigid foot was equal to the deepest part of
321 the real moving foot at mid-stance. Timing was such that the indentation and removal of the rigid
322 foot took the same number of frames as the experimental trial being simulated, i.e. the simulated
323 time taken to 'stamp' the rigid foot was equal to the real timing of the original footstep. This
324 most simplistic scenario was followed first by a single rigid foot rotating to approximate a heel-

325 toe cycle, and then by a two-part foot in which the toes were able to rotate as an object
326 independently of the foot (i.e., with a simple hinge at the approximate positions of the
327 metatarsophalangeal joints). The single rotating foot object was animated to sink in the substrate
328 such that the maximum depth of the metatarsal heads matched the depth of the metatarsal heads
329 in the biplanar X-ray data. While this meant the majority of the foot approximated the motion of
330 the bi-planar X-ray data, the toes necessarily sank much farther due to significant rotation. The
331 two-part model alleviated this by allowing the toes to remain more horizontal as the heel lifted
332 off the substrate. This two-part rigid body simulation is analogous to previous footprint
333 simulation work (22,28) in which individual toe segments were treated as separate translating
334 and rotating rigid bodies.

335 However, these rigid-body models failed to capture subtle deformations of the human
336 foot, particularly involving flexibility of the arches. Our final simulation used the animated high-
337 resolution foot mesh directly, capturing as much of the reconstructed motion as possible. To do
338 this, mesh face and vertex positions were output at a far greater temporal-resolution; 1000 frames
339 per second. LIGGGHTs input files ran 1000 timesteps (each of 0.000001 seconds real time)
340 between each frame to translate the mesh from one position to the next. This produced the most
341 ‘realistic’ simulations, incorporating all motion of the deforming foot as derived from the skin
342 markers placed on the subject. Simulations were visualized using OVITO (v. 3.0.0) (33).

343

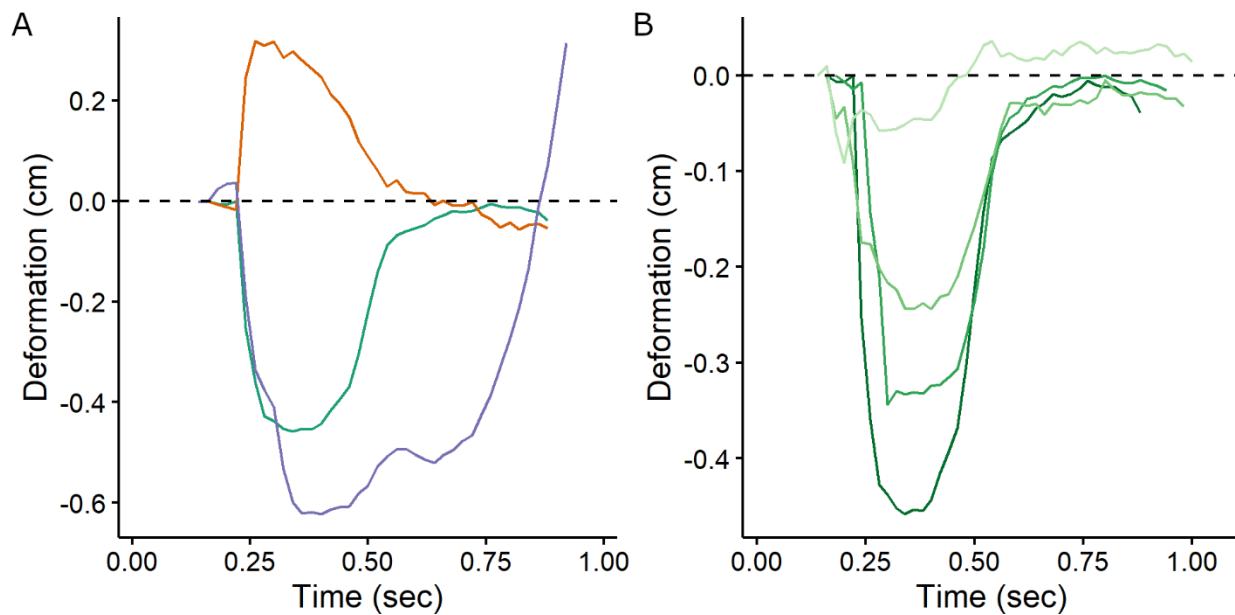
344 **Results and Discussion**

345 Using the methods described above, we successfully built data-driven 3-D animations of
346 deformable feet navigating deformable substrates to produce tracks (Supplementary Video S1).
347 Since the methodological developments are the focus of this paper, we present data from a single

348 subject as a case study to demonstrate the variety of analyses that are permitted through the
349 application of these novel methods.

350 The first area in which we can apply these techniques is to study 3-D kinematics of the
351 foot at the substrate interface. The biplanar X-ray technique presented here provides a window
352 for direct visualization of the foot-substrate interface while a human foot travels into, and
353 interacts with, both rigid and deformable substrates. As in previous studies (25), the 3-D
354 positions of external foot markers, visualized through biplanar X-ray, can be used to quantify 3-
355 D deformations of the plantar surface of the foot during its interactions with these various
356 substrates. For example, continuous measurements of heel compression, heel expansion, and
357 longitudinal arch deformation can be collected throughout the duration of stance phase to
358 understand soft tissue behavior in these regions of the foot (Fig. 5).

359



360

361 **Figure 5.** 3-D deformation of the foot of one individual walking across multiple substrates. A)
362 Continuous measurements of heel height (green), heel width (orange), and medial longitudinal
363 arch height (purple) during one trial on carbon fiber. Each measurement is zeroed based on its

364 first possible measurement (prior to initial contact, when the foot first entered both biplanar X-
365 ray video frames). B) Sample plots showing deformation of the heel (change in vertical height)
366 in one subject walking across four different substrates. Substrates become more deformable as
367 they transition from darker to lighter shades of green (carbon fiber is the darkest green, “firm”
368 mud is the second darkest, “hydrated 2.5” mud is the second lightest, and “hydrated 5” mud is
369 the lightest).

370

371 Figure 5 portrays temporal and substrate-driven patterns of foot deformation consistent
372 with those previously observed by Hatala et al. (25). The external surface of the heel
373 simultaneously compressed vertically and expanded horizontally as the calcaneal fat pad
374 dissipated impact forces (Fig. 5A), a pattern which has been well-studied experimentally (34–
375 36). The medial longitudinal arch initially flattened as the foot was loaded, but at terminal stance
376 phase it eventually reached a height that exceeded its initial, unloaded, state (Fig. 5A), consistent
377 with results from other experimental studies of longitudinal arch function (37). Comparisons
378 across substrates likewise followed patterns observed previously by Hatala et al. (25). For
379 example, the heel compressed to greater degrees as subjects walked over more rigid substrates
380 (Fig. 5B). Clearly these are not the only types of dynamic measurements that can be acquired,
381 and a variety of 3-D kinematic studies would be possible through this approach. We simply
382 emphasize here that our experimental protocol offers several directions to study foot-substrate
383 interactions across rigid and deformable substrates using external marker-based kinematics.

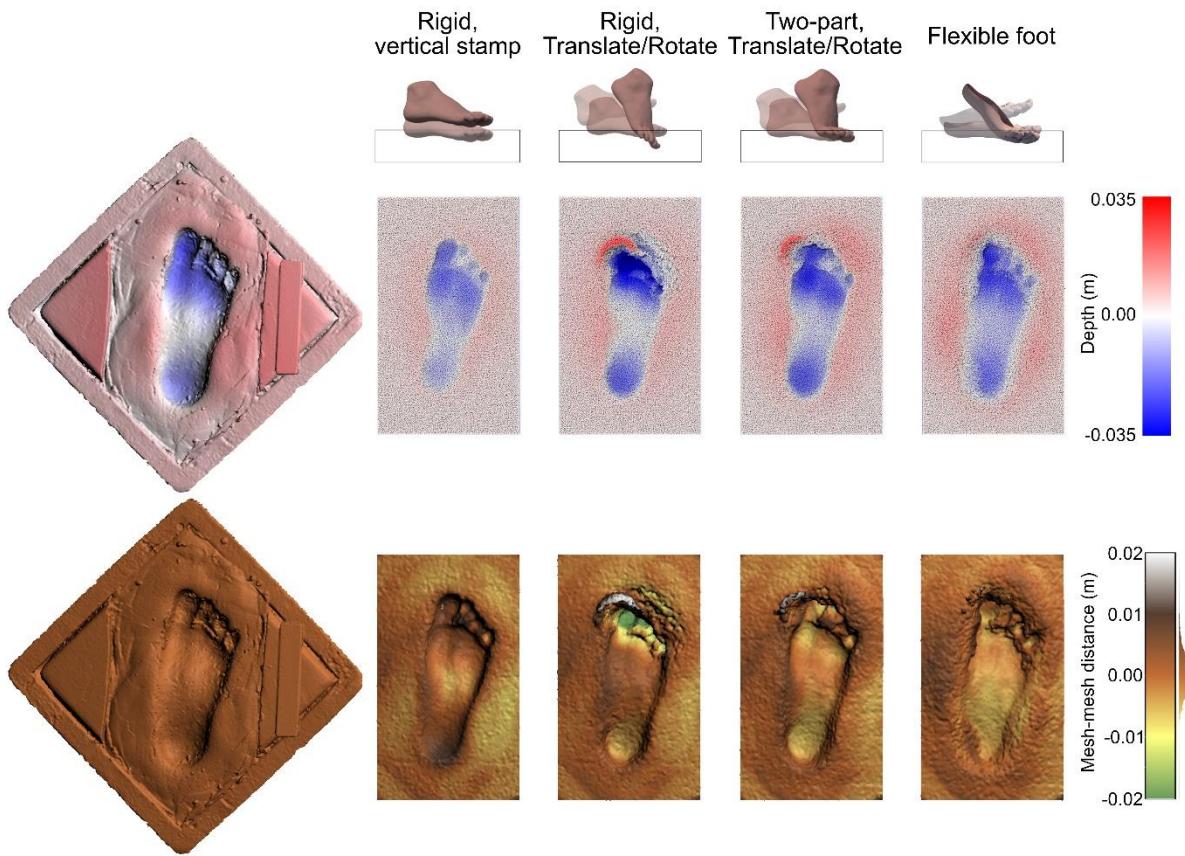
384 Building upon studies of pure foot deformation and motion, the integration of high-
385 resolution 3-D models of both feet and tracks within the same animation scene provides
386 opportunities to observe directly the extent and nature of correspondence between external foot

387 motions and the morphology of the final track that was produced. Previous studies have
388 highlighted the lack of direct correspondence between foot motion and track morphology (25)
389 and similar patterns were observed here. It is evident that final track morphology is not simply a
390 Boolean-type subtraction of the foot's trajectory through the substrate. While the lack of
391 correspondence between foot trajectories and final track morphology can be observed from the
392 results of 3-D animations of experimental trials, a true understanding of these differences
393 requires knowledge of human track ontogeny. Such knowledge can be gained through track
394 simulations, which allow one to visualize and understand the patterns of substrate flow that
395 generate specific aspects of track morphology. Here we explored as a case study a single trial
396 from our biplanar X-ray experiments, in which a subject walked across "hydrated 5" mud to
397 produce a track. The 3-D scan of that track was directly compared with simulated tracks that
398 were produced following the track simulation protocols described above.

399 By iteratively increasing the complexity of the deformation and motion of the animated
400 foot, we achieved simulations that eventually produced track morphologies that closely matched
401 those produced in biplanar X-ray experiments (Fig. 6, Table 1). The simplest simulation, in
402 which a rigid foot model vertically stamped a substrate, actually generated a track morphology
403 with the smallest average pairwise distance from the 3-D scanned track (Table 1) and that looked
404 qualitatively realistic. However, the similarities between the simulated and scanned tracks were
405 largely confined to the region of the forefoot (Fig. 6). This was unsurprising, since the simulated
406 foot trajectories were configured such that maximum depth beneath the metatarsal heads
407 matched the depths to which the metatarsal heads were observed to travel in biplanar X-ray
408 experiments (i.e., all simulations are most likely to match the 3-D scanned track in the region of
409 the forefoot). The "vertical stamp" produced a track that was noticeably shallower and narrower

410 than the scanned track in the region of the heel, and that had an overall less longitudinally arched
411 shape. This track also lacked the displacement rims that surrounded the perimeter of the scanned
412 track.

413



414

415 **Figure 6.** Direct comparisons between 3-D scan of track from biplanar X-ray experiments (left)
416 and 3-D meshes of tracks produced in various particle simulations (right). Simulations increase
417 in complexity from left to right, from a vertical stamp of a rigid foot to a step taken by a fully
418 flexible foot, whose motions and deformations were driven by real data from biplanar X-ray
419 experiments. Top row shows track depths (in meters) as measured from the ground plane.
420 Bottom row shows pairwise distances between each simulated track and the actual 3-D scanned
421 track. Differences between simulation conditions are subtle, but overall the most complicated

422 animation/simulation converges on a track morphology that is most similar to the one actually
423 produced in biplanar X-ray experiments.

424

425 **Table 1.** Summary statistics for pairwise distance comparisons between simulated tracks and 3-
426 D scanned track from biplanar X-ray experiments.

| Simulation type | Mean distance (cm) | Standard deviation (cm) |
|---------------------------------|--------------------|-------------------------|
| Rigid foot, vertical stamp | 0.0062 | 0.3446 |
| Rigid foot, translate/rotate | -0.0286 | 0.5980 |
| Two-part foot, translate/rotate | 0.0556 | 0.3511 |
| Fully flexible animated foot | 0.0176 | 0.2885 |

427

428 By adding motion to the rigid foot model (translating and rotating a rigid foot), we
429 produced simulated tracks that had greater relative elevation beneath the longitudinal arch but
430 that were otherwise quite different from the 3-D scanned track. Toe impressions were extremely
431 deep, the heel impression was deeper than observed in the scanned track, and a very noticeable
432 extrusion feature was generated at the tip of the hallux (Fig. 6). Displacement rims were still not
433 as prominent as they were in the 3-D scanned experimental track. Adding a simple hinge to
434 convert the rigid foot into a two-part model (allowing the foot to deform at the approximate
435 positions of the metatarsophalangeal joints) remedied some but not all of these inaccuracies.
436 Forefoot (including toe) impressions were overall more similar to those of the 3-D scanned track,

437 but the heel impression was still deeper and the extrusion feature at the tip of the hallux was still
438 generated (Fig. 6).

439 Implementing a fully mobile and deformable foot animation led to simulated tracks that
440 most closely matched those observed in biplanar X-ray experiments. The mean distance between
441 the simulated and 3-D scanned tracks was only second lowest but the standard deviation was the
442 smallest, indicating that this simulation varied the least of the four scenarios from the original
443 scanned surface (Table 1). The simulated track was similar in relative depths across the forefoot
444 (including toe) impressions, relative depths in the region of the heel, and in the pattern of the
445 displacement rim surrounding the perimeter of the track (Fig. 6). It was also the widest track in
446 the mid-foot, which matched most closely with the real track. The simulated track had a slightly
447 deeper impression beneath the longitudinal arch than did the 3-D scanned track, but this
448 difference was relatively subtle.

449 It is clear from our simulated tracks that, as might be expected, incorporation of more
450 complex motions and soft-tissue deformations results in a more true-to-life final track
451 morphology. That the real track differed substantially from the ‘stamp’ simulation demonstrates
452 once again that “footprints are not feet” and should not be interpreted as direct reflections of
453 plantar foot anatomy (29). Our simulated tracks also highlight caution in using simple metrics
454 such as mean mesh-mesh distances to compare tracks; the complex 3D topography means that
455 mean distances can be low, even when tracks are clearly qualitatively different.

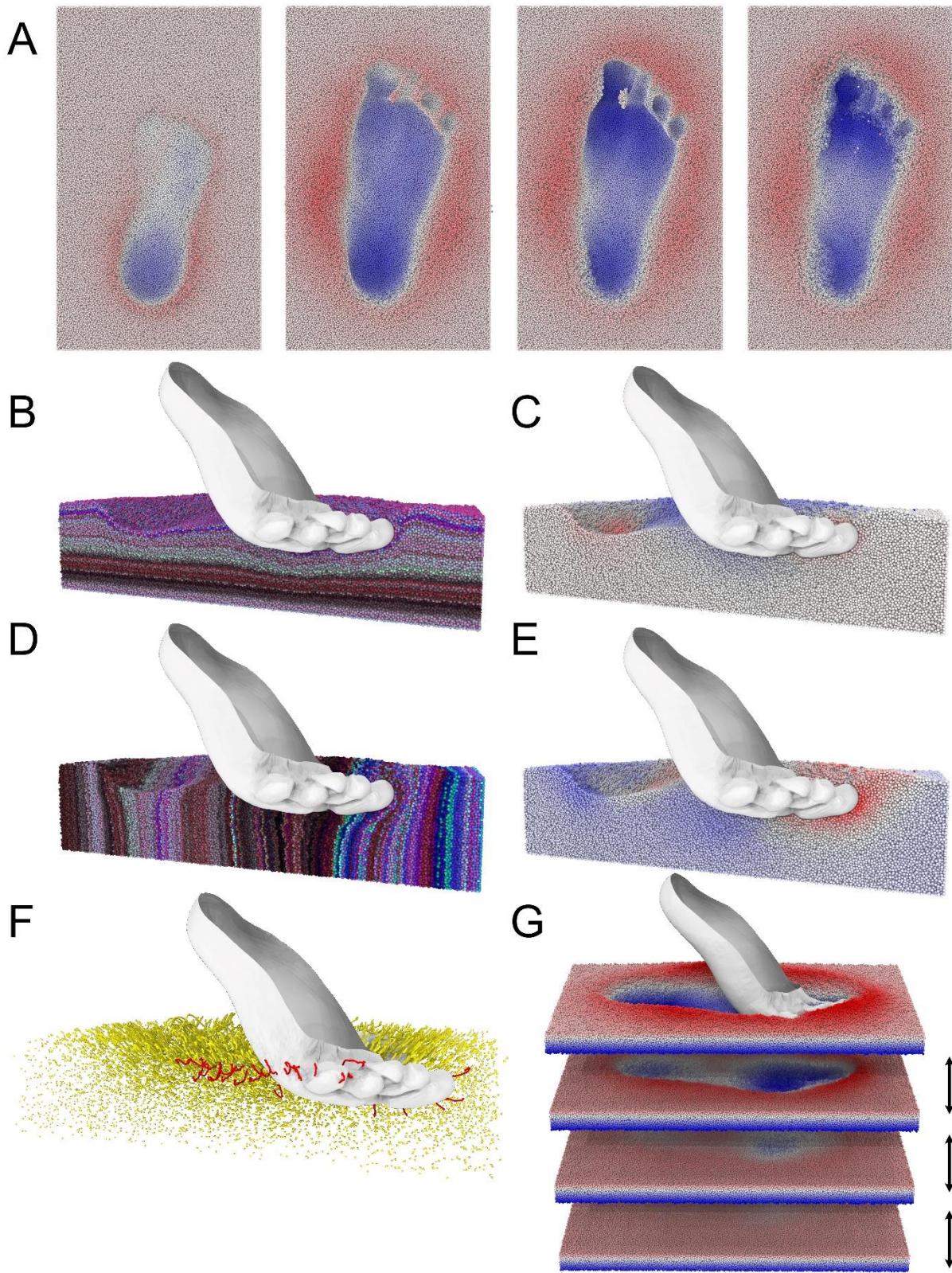
456 Focusing on our most complex simulation (deformable foot), the qualitative and
457 quantitative similarity between simulated track and real scanned track is gratifying, and indicates
458 that the real motions of the foot and substrate are captured by our workflow. Minor differences
459 between the final simulated track and the 3D-scan of the real impression can be attributed to

460 simulation parameters, particularly particle size and cohesion, though refining these parameters
461 further would require substantial iterative simulations, which for the purposes of this study were
462 deemed unnecessary. The nature of how the sediments are mixed and set-up during the
463 experimental protocol means that the bulk properties of the experimental substrate (particularly
464 as it overlies elastically-behaving foam) would be difficult to ascertain from a smaller, and thus
465 easier to simulate, sample. As such, we base our input parameters on what makes the output most
466 like the scanned track, but as elaborated on previously (28) significant deviations between
467 simulation and reality would indicate our input parameters are incorrect. We therefore consider
468 our simulation, based on its qualitative and quantitative similarity to the scanned track, to
469 accurately represent the pattern of surface and sub-surface substrate deformation that occurred
470 during the biplanar X-ray experiment.

471 Armed with this complete simulation of animated, deforming foot morphology and a
472 deformable substrate responding to that foot, we are able to visualize and explore the formation
473 of the track - its ontogeny - in a multitude of ways at and beneath the sediment surface (Fig. 7).
474 Examining a sequence of time steps during the foot-substrate interaction allows us to visualize
475 the temporal process of track development (Fig. 7A). Using randomized bands of colour oriented
476 either vertically or horizontally, enables visualization of the directions and magnitudes of particle
477 motion within the substrate (Figs. 7B and 7D). Color gradients can also be applied to individual
478 particles, in order to visualize how far they move in various directions (Figs. 7C and 7E). Particle
479 trajectories can be traced in order to track motions of individual particles or groups of particles
480 within the substrate throughout the track forming process (Fig. 7F). For instance, selecting
481 particles in the displacement rims and generating trajectories backwards, we can identify where
482 the raised sediment has been pushed from. Subsurface layers can be exposed, presenting

483 transmitted undertracks (Fig. 7G). Ultimately there are countless directions that one can pursue
484 to visualize track ontogeny, and understand how various aspects of track morphology were
485 generated. We do not exhaustively list the possibilities here, but merely emphasize a variety of
486 visualization techniques that can reveal previously hidden aspects of the track formation process.

487



489 **Figure 7.** Examples of visualization methods applicable to our simulated tracks. A) Track
490 ontogenetic sequence at ~25, 50, 75, and 100% of stance phase. Colour scale indicates height,
491 and difference between darkest blue-red is 7 cm. B) Randomized horizontal colouring, exposed
492 through longitudinal section, provides a view comparable with observing a laminated sediment.
493 C) Medio-lateral motion of individual particles can be represented with colour, blue particles
494 having moved medially, and red particles having moved laterally. D) and E) Visualize
495 forward/backward motion of particles as either randomized vertical colouration (D) or colour-
496 coded such that red indicates forward motion, blue indicates backward motion (E). F)
497 Demonstrates particle vectors throughout the track forming process. Particles of interest, such as
498 those in red which form the displacement rims, can be tracked separately and individually. G)
499 The simulated track can be split at virtual bedding planes, exposing a sequence of penetrative
500 and transmitted undertracks.

501

502 **Conclusions**

503 The combination of biplanar X-ray, 3-D animation, and particle simulation methods that
504 we have introduced and applied here have the potential to inform a wide variety of research
505 questions related to how locomotion varies across substrates with different mechanical
506 properties, and how tracks can record those variations. Instruments that are ubiquitous to
507 biomechanics labs, such as force plates, pressure pads, and optical motion capture systems,
508 provide richly detailed understandings of how our feet function during locomotion. However,
509 force- and pressure-sensing instruments are typically rigid and the opacity of feet and substrates
510 conceal the interactions that occur at the foot-substrate interface, so these instruments are for the
511 most part limited to studying locomotion on rigid surfaces. The hidden interactions between foot

512 and deformable substrate are of interest to researchers across many disciplines that seek to better
513 understand their mechanics. For example, in biorobotics, a great deal of attention has been
514 devoted to understanding how animals traverse irregular, deformable terrain. It has been
515 challenging to build robots that can navigate natural environments and their inherent
516 unpredictability, in part due to limited abilities to observe and measure mechanical interactions at
517 the foot-substrate interface (38,39). In human biomechanics, understandings of locomotion and
518 foot function across non-rigid substrates are similarly limited. It is known that humans alter their
519 kinematics on deformable substrates, and that the energetic costs of locomotion increase with
520 substrate compliance (40–42). However, it has been exceedingly difficult to observe and quantify
521 the manners in which human feet engage with non-rigid substrates. The methods described here
522 are transferable to these and other systems, and have the potential to open windows on
523 previously unobservable biomechanical phenomena. This emphasizes the interdisciplinarity that
524 is inherent to these approaches.

525 Within paleoanthropology, the methods developed here substantially expand the toolkit
526 that can be applied to analyze hominin tracks. Previous experimental studies, including our own,
527 have relied on the comparative method to determine whether and how various hominin tracks
528 differ from each other, and to develop anatomical and/or functional hypotheses for those
529 differences (9,11,43–48). The methods presented here focus instead on building knowledge of
530 human track ontogeny, in order to understand how particular anatomical or functional patterns
531 lead to the development of specific track morphologies. Through validated track simulation
532 methods, the combinations of foot anatomy and motion that would be capable of producing
533 particular fossil track morphologies can be reverse-engineered (28). When synthesized with
534 “functional” analyses of skeletal fossils (e.g., analyses of trabecular bone, cross-sectional

535 geometry, and/or articular morphology), these simulation-based analyses of fossil hominin tracks
536 provide an unparalleled route to explicitly test and develop hypotheses regarding fossil hominin
537 locomotion.

538

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549

550 **References**

- 551 1. Darwin C. *The descent of man, and selection in relation to sex*. London: J. Murray;
552 1871. 898 p.
- 553 2. DeSilva J, McNutt E, Benoit J, Zipfel B. One small step: A review of Plio-
554 Pleistocene hominin foot evolution. *Am J Phys Anthropol.* 2019;168:63–140.
- 555 3. Day MH, Napier JR. Fossil foot bones. *Nature.* 1964;201(4923):969–70.
- 556 4. Jungers WL, Harcourt-Smith WEH, Wunderlich RE, Tocheri MW, Larson SG,
557 Sutikna T, et al. The foot of *Homo floresiensis*. *Nature.* 2009;459(7243):81–4.

- 558 5. Harcourt-Smith WEH, Throckmorton Z, Congdon KA, Zipfel B, Deane AS,
559 Drapeau MSM, et al. The foot of *Homo naledi*. *Nat Commun.* 2015;6:1–8.
- 560 6. DeSilva JM, Gill CM, Prang TC, Bredella MA, Alemseged Z. A nearly complete
561 foot from Dikika, Ethiopia and its implications for the ontogeny and function of
562 *Australopithecus afarensis*. *Sci Adv.* 2018;4(7):eaar7723.
- 563 7. Gatesy SM, Middleton KM, Jenkins Jr. FA, Shubin NH. Three-dimensional
564 preservation of foot movements in Triassic theropod dinosaurs. *Nature.*
565 1999;399:141–4.
- 566 8. Manafzadeh AR, Padian K. ROM mapping of ligamentous constraints on avian hip
567 mobility: Implications for extinct ornithodirans. *Proc R Soc B Biol Sci.*
568 2018;285(1879).
- 569 9. Bennett MR, Harris JWK, Richmond BG, Braun DR, Mbua E, Kiura P, et al. Early
570 hominin foot morphology based on 1.5-million-year-old footprints from Ileret,
571 Kenya. *Science.* 2009;323(5918):1197–201.
- 572 10. Ashton N, Lewis SG, De Groote I, Duffy SM, Bates M, Bates R, et al. Hominin
573 footprints from early Pleistocene deposits at Happisburgh, UK. *PLoS One.*
574 2014;9(2):e88329.
- 575 11. Hatala KG, Roach NT, Ostrofsky KR, Wunderlich RE, Dingwall HL, Villmoare
576 BA, et al. Footprints reveal direct evidence of group behavior and locomotion in
577 *Homo erectus*. *Sci Rep.* 2016;6:28766.
- 578 12. Hatala KG, Roach NT, Ostrofsky KR, Wunderlich RE, Dingwall HL, Villmoare
579 BA, et al. Hominin track assemblages from Okote Member deposits near Ileret,

- 580 Kenya, and their implications for understanding fossil hominin paleobiology at 1.5
581 Ma. *J Hum Evol.* 2017;112:93–104.
- 582 13. Masao FT, Ichumbaki EB, Cherin M, Barili A, Boschian G, Iurino DA, et al. New
583 footprints from Laetoli (Tanzania) provide evidence for marked body size variation
584 in early hominins. *eLife.* 2016;5:29.
- 585 14. Altamura F, Bennett MR, D'Août K, Gaudzinski-Windheuser S, Melis RT,
586 Reynolds SC, et al. Archaeology and ichnology at Gombore II-2, Melka Kunture,
587 Ethiopia: everyday life of a mixed-age hominin group 700,000 years ago. *Sci Rep.*
588 2018;8(1):1–11.
- 589 15. Altamura F, Bennett MR, Marchetti L, Melis RT, Reynolds SC, Mussi M.
590 Ichnological and archaeological evidence from Gombore II OAM, Melka Kunture,
591 Ethiopia: An integrated approach to reconstruct local environments and biological
592 presences between 1.2 and 0.85 Ma. *Quat Sci Rev.* 2020;244:106506.
- 593 16. Bennett MR, Falkingham P, Morse SA, Bates K, Crompton RH. Preserving the
594 impossible: Conservation of soft-sediment hominin footprint sites and strategies for
595 three-dimensional digital data capture. *PLoS One.* 2013;8(4).
- 596 17. Falkingham PL, Bates KT, Avanzini M, Bennett MR, Bordy EM, Breithaupt BH, et
597 al. A standard protocol for documenting modern and fossil ichnological data.
598 *Palaeontology.* 2018;
- 599 18. Minter NJ, Braddy SJ, Davis RB. Between a rock and a hard place: Arthropod
600 trackways and ichnotaxonomy. *Lethaia.* 2007;40(4):365–75.
- 601 19. Padian K, Olsen PE. The fossil trackway *Pteraichnus*: not pterosaurian, but
602 crocodilian. *J Paleontol.* 1984;178–84.

- 603 20. Falkingham PL. Interpreting ecology and behaviour from the vertebrate fossil track
604 record. *J Zool.* 2014;292(4):222–8.
- 605 21. Morse SA, Bennett MR, Liutkus-Pierce C, Thackeray JF, McClymont J, Savage R,
606 et al. Holocene footprints in Namibia: the influence of substrate on footprint
607 variability. *Am J Phys Anthropol.* 2013;151(2):265–79.
- 608 22. Falkingham PL, Gatesy SM. The birth of a dinosaur footprint: Subsurface 3D
609 motion reconstruction and discrete element simulation reveal track ontogeny. *Proc
610 Natl Acad Sci.* 2014;111(51):18279–84.
- 611 23. Maladen R, Ding Y, Li C, Goldman DI. Undulatory swimming in sand: Subsurface
612 locomotion of the sandfish lizard. *Science.* 2009;325(5938):314–8.
- 613 24. Ellis RG, Gatesy SM. A biplanar X-ray method for three-dimensional analysis of
614 track formation. *Palaeontol Electron.* 2013;16(1):1T.
- 615 25. Hatala KG, Perry DA, Gatesy SM. A biplanar X-ray approach for studying the 3D
616 dynamics of human track formation. *J Hum Evol.* 2018;121:104–18.
- 617 26. Cundall PA, Strack ODL. A discrete numerical model for granular assemblies.
618 *Géotechnique.* 1979;29:47–65.
- 619 27. Kloss C, Goniva C. LIGGGHTS - Open source discrete element simulations of
620 granular materials based on Lammps. In: TMS, editor. *Supplemental Proceedings:
621 Materials Fabrication, Properties, Characterization, and Modeling, Volume 2.*
622 Hoboken: John Wiley & Sons, Inc; 2011. p. 781–8.
- 623 28. Falkingham PL, Turner ML, Gatesy SM. Constructing and testing hypotheses of
624 dinosaur foot motions from fossil tracks using digitization and simulation.
625 *Palaeontology.* 2020;1–16.

- 626 29. Gatesy SM, Falkingham PL. Neither bones nor feet: track morphological variation
627 and ‘preservation quality.’ *J Vertebr Paleontol.* 2017;e1314298.
- 628 30. Brainerd EL, Baier DB, Gatesy SM, Hedrick TL, Metzger KA, Gilbert SL, et al. X-
629 ray reconstruction of moving morphology (XROMM): precision, accuracy and
630 applications in comparative biomechanics research. *J Exp Zool A.*
631 2010;313(5):262–79.
- 632 31. Knörlein BJ, Baier DB, Gatesy SM, Laurence-Chasen JD, Brainerd EL. Validation
633 of XMALab software for marker-based XROMM. *J Exp Biol.* 2016;219(23):3701–
634 11.
- 635 32. Baier DB. XROMM Maya Tools. 2018. Available from:
636 https://bitbucket.org/xromm/xromm_mayatools/src/master/
- 637 33. Stukowski A. Visualization and analysis of atomistic simulation data with OVITO -
638 the Open Visualization Tool. *Modelling Simul Mater Sci Eng.* 2010;18:015012.
- 639 34. De Clercq D, Aerts P, Kunnen M. The mechanical characteristics of the human heel
640 pad during foot strike in running: An in vivo cineradiographic study. *J Biomech.*
641 1994;27(10):1213–22.
- 642 35. Gefen A, Megido-Ravid M, Itzhak Y. In vivo biomechanical behavior of the
643 human heel pad during the stance phase of gait. *J Biomech.* 2001;34:1661–1665.
- 644 36. Chi KJ, Schmitt D. Mechanical energy and effective foot mass during impact
645 loading of walking and running. *J Biomech.* 2005;38(7):1387–95.
- 646 37. Caravaggi P, Pataky T, Gunther M, Savage R, Crompton R. Dynamics of
647 longitudinal arch support in relation to walking speed: contribution of the plantar
648 aponeurosis. *J Anat.* 2010;217(3):254–61.

- 649 38. Ijspeert AJ. Biorobotics: using robots to emulate and investigate agile locomotion.
- 650 Science. 2014;346(6206):196–203.
- 651 39. Aguilar J, Zhang T, Qian F, Kingsbury M, McInroe B, Mazouchova N, et al. A
- 652 review on locomotion robophysics: The study of movement at the intersection of
- 653 robotics, soft matter and dynamical systems. Reports Prog Phys.
- 654 2016;79(11):110001.
- 655 40. Ferris DP, Louie M, Farley CT. Running in the real world: Adjusting leg stiffness
- 656 for different surfaces. Proc R Soc B Biol Sci. 1998;265(1400):989–94.
- 657 41. Lejeune TM, Willems PA, Heglund NC. Mechanics and energetics of human
- 658 locomotion on sand. J Exp Biol. 1998;201:2071–80.
- 659 42. Kerdok AE, Biewener AA, McMahon TA, Weyand PG, Herr HM. Energetics and
- 660 mechanics of human running on surfaces of different stiffnesses. J Appl Physiol.
- 661 2002;92(2):469–78.
- 662 43. Day MH, Wickens EH. Laetoli Pliocene hominid footprints and bipedalism. Nature.
- 663 1980;286:385–7.
- 664 44. Tuttle RH, Webb DM, Weidl E, Baksh M. Further progress on the Laetoli trails. J
- 665 Archaeol Sci. 1990;17:347–62.
- 666 45. Meldrum DJ. Midfoot Flexibility, Fossil Footprints, and Sasquatch Steps: New
- 667 Perspectives on the Evolution of Bipedalism. J Sci Explor. 2004;18(1):65–79.
- 668 46. Raichlen DA, Gordon AD, Harcourt-Smith WEH, Foster AD, Haas WR. Laetoli
- 669 footprints preserve earliest direct evidence of human-like bipedal biomechanics.
- 670 PLoS One. 2010;5(3):e9769.

- 671 47. Crompton RH, Pataky TC, Savage R, D'Août K, Bennett MR, Day MH, et al.
- 672 Human-like external function of the foot, and fully upright gait, confirmed in the
- 673 3.66 million year old Laetoli hominin footprints by topographic statistics,
- 674 experimental footprint-formation and computer simulation. *J R Soc Interface*.
- 675 2012;9(69):707–19.
- 676 48. Hatala KG, Demes B, Richmond BG. Laetoli footprints reveal bipedal gait
- 677 biomechanics different from those of modern humans and chimpanzees. *Proc R Soc*
- 678 B. 2016;283:20160235.