Global Quantitative SILAC Phosphoproteomics Reveals Differential Phosphorylation Is Widespread between the Procyclic and Bloodstream Form Lifecycle Stages of *Trypanosoma brucei*

Michael D. Urbaniak, David M. A. Martin, and Michael A. J. Ferguson*

Division of Biological Chemistry and Drug Discovery, College of Life Sciences, University of Dundee, Dundee DD1 5EH, United Kingdom

Supporting Information

**ABSTRACT:** We report a global quantitative phosphoproteomic study of bloodstream and procyclic form *Trypanosoma brucei* using SILAC labeling of each lifecycle stage. Phosphopeptide enrichment by SCX and TiO2 led to the identification of a total of 10096 phosphorylation sites on 2551 protein groups and quantified the ratios of 8275 phosphorylation sites between the two lifecycle stages. More than 9300 of these sites (92%) have not previously been reported. Model-based gene enrichment analysis identified over representation of Gene Ontology terms relating to the flagella, protein kinase activity, and the regulation of gene expression. The quantitative data reveal that differential protein phosphorylation is widespread between bloodstream and procyclic form trypanosomes, with significant intraprotein differential phosphorylation. Despite a lack of dedicated tyrosine kinases, 234 phosphotyrosine residues were identified, and these were 3–4 fold over-represented among site changing >10-fold between the two lifecycle stages. A significant proportion of the *T. brucei* kinome was phosphorylated, with evidence that MAPK pathways are functional in both lifecycle stages. Regulation of gene expression in *T. brucei* is exclusively post-transcriptional, and the extensive phosphorylation of RNA binding proteins observed may be relevant to the control of mRNA stability in this organism.

**KEYWORDS:** phosphorylation, SILAC, Trypanosoma brucei, quantitative proteomics, phosphoproteomics

**INTRODUCTION**

The protozoan parasite *Trypanosoma brucei* is the etiological agent of African sleeping sickness, also known as Human African Trypanosomiasis (HAT), which is transmitted by the bite of an infected tsetse fly. The disease is usually fatal if left untreated and is estimated to be responsible for around 10000 deaths per annum in sub-Saharan Africa. Current treatments are expensive, toxic and difficult to administer, leaving an urgent unmet need for new therapeutic agents. *T. brucei* has a complex digenetic lifecycle between an insect vector and a mammalian host, and its ability to adapt its proteome to these disparate environments is essential to its survival and virulence. During the early stages of an infection, the clinically relevant bloodstream form of the parasite proliferates in the blood and lymph of the human host, then in the second stage enters the cerebral-spinal fluid and brain, resulting in coma and death. When a tsetse fly feeds on an infected host, the parasite is ingested with the blood meal, triggering parasite differentiation into the procyclic form to enable survival in its new environment.

Both the procyclic form and bloodstream form of the parasite may be cultured *in vitro*. We have recently quantified the changes in protein expression between the procyclic and bloodstream form lifecycle stages using SILAC labeling of procyclic cells, demonstrating that about 10% of the observed proteome is differentially regulated by more than 5-fold. The fold-change in protein level showed a strong correlation (Pearson correlation 0.86) to changes in mRNA level recorded by transcriptomic studies, despite the atypical mechanism of gene expression in trypanosomes, where genes are transcribed in large polycistronic units that are trans-spliced into mature mRNAs for translation, with mRNA abundance controlled by stability.

There is evidence that protein phosphorylation may be important in *T. brucei*. Nett et al. used sequential strong-cation exchange (SCX) and TiO2 enrichment of phosphopeptides derived from the cytosolic fraction of bloodstream form parasites to identify 1204 phosphorylation sites on 491 proteins, and immunoprecipitation with antiphosphotyrosine antibodies to identify 34 phosphotyrosine sites induced in response to the treatment of procyclic form cells with hydrogen peroxide. However, to date, there has been no quantitative comparison of the phosphoproteomes of the two lifecycle stages. We set out to define the global protein phosphorylation status of the two lifecycle stages to explore the biological processes that might be modulated by phosphorylation, and to highlight stage-specific
phosphoproteomic changes. Here, we report a global quantitative phosphoproteomic study of bloodstream and procyclic form *Trypanosoma brucei* using SILAC labeling of each lifecycles stage that identifies 9314 previously unidentified phosphorylation sites. The results show that differential phosphorylation is widespread between the procyclic and bloodstream form and that it adds significant complexity to the changes in the proteome.

**EXPERIMENTAL SECTION**

**Cell Culture Media**

**HM19-T**, a modification of the original IMDM-based HMI-9, uses 56 μM 1-thioglycerol in place of 200 μM 2-mercaptoethanol, 2 mM glutamax I (Invitrogen) in place of 4 mM L-glutamine, and contains 10% heat-inactivated fetal bovine serum (PAA). Similarly, HMI11-T is a modification of the original HMI11 (HMI-9 lacking serum plus) that use 56 μM 1-thioglycerol and 2 mM glutamax I with 10% heat-inactivated fetal bovine serum.

HMI11-SILAC — RK was prepared in the same way as HMI11-T but using IMDM depleted of L-Arginine, L-Lysine (Sigma) and 10% dialyzed heat-inactivated fetal bovine serum (10 kDa molecular weight cutoff, PAA) and was supplemented with 4 mg/mL folic acid. The HMI11-SILAC — RK was supplemented with either normal L-Arginine and L-Lysine (HMI11-SILAC + R6K6) or with L-Arginine U-13C6/L-Lysine-2H4 (HMI11-SILAC + R6K6), Cambridge Isotope Laboratories) at 30% of the original HMI11 concentration (120 μM) respectively unless otherwise stated. Original SDM-79 medium and SILAC SDM-79 medium (SDM-79 + R6K6) were prepared as described previously.

**Cell Culture**

Procyclic form *Trypanosoma brucei* Lister 427 clone 29.13.6 cells were grown at 28 °C without CO2 in SDM-79 in the presence of 15 μg/mL G418 at 37 °C in a 5% CO2 incubator. Both cell lines have been genetically modified to express the T7 RNA polymerase and the tetracycline repressor protein, as described by Wirtz et al. Cell dimensions were measured using a CASY cell counter.

For the growth curves, the *T. brucei* bloodstream form cells were washed twice in 10 mL HMI11-SILAC + R6K6 for 11–12 cell divisions and hypotonically lysed as described above. To reduce the sample complexity the proteins were fractionated by SDS-PAGE, and a band corresponding to 25–50 kDa molecular weight range was excised and subjected to in-gel tryptic digest prior to analysis by LC—MS/MS. Ten peptides were chosen at random and the relative abundance of the major isotopic peak of heavy (arginine-13C6/lysine-2H4) and light (arginine-12C6/lysine-2H4) forms were measured using the extracted ion chromatogram function in Excalibur (Thermo Scientific).

**Estimating Efficiency of SILAC Labeling**

Bloodstream form cells were grown in HMI11-SILAC + R6K6 for >14 days prior to inoculation of female Balb/c mice (three per condition) by intraperitoneal injection of 1 × 10^5, 1 × 10^4 or 1 × 10^3 cells in the same media. Daily tail bleeds were used to monitor the number of parasites in the bloodstream using a haemocytometer, and animals were euthanized when the parasite burden exceeded 5 × 10^6 cells/mL prior to overt distress.

**Comparative Proteomic Analysis**

A 1:1 mix of procyclic form cells grown in heavy-SILAC media and bloodstream form cells grown in heavy-SILAC media was solubilized with SDS and tryptic peptides were generated by FASP. Peptides were fractionated by SCX prior to analysis by liquid chromatography tandem mass spectrometry as described previously, except only 5 SCX fractions were analyzed and technical duplicates were not performed. The previously reported the analysis of a 1:1 mix of procyclic form cells grown in heavy-SILAC media and bloodstream form cells grown in light-SILAC media, consisting of technical duplicates of eight SCX fractions and ten SDS-PAGE fractions, was reprocessed to ensure accurate comparison with the current data.

**Filter Aided Sample Preparation**

Peptide samples for analysis by mass spectrometry were prepared by modification of the filter-aided sample preparation procedure (FASP). Briefly, samples were desalted and combined according to the experiment design to give a total of 2.5 × 10^9 lysed cells (0.5 mL), solubilized with 4% SDS and then reductively alkylated in a 30000 molecular weight cutoff vertical spin filtration unit (Vivascience) using the FASP procedure adapted for the larger volumes used here. The sample was digested with 1:100 ratio (w/w) of trypsin gold (Promega) in the filtration unit for 18 h at 37 °C, tryptic peptides were eluted by centrifugation, and the filter washed sequentially with 1 mL of 50 mM NH4HCO3 and 1 mL of 0.5 M NaCl. The combined eluent was desalted using a 500 mg C18 cartridge (SepPak, Waters) and lyophilized.

**Strong cation Exchange Chromatography**

Strong cation exchange (SCX) was performed on an Agilent 1120 compact LC using a 3.0 × 200 mm 5 μm polysulfolyl aspartamide column (Poly LC) with a flow rate of 350 μL/min.
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and detection at 220 nm. Phosphopeptides were enriched using the method of Beausoleil et al.12 Dried peptides were dissolved in 200 μL of solvent A (5 mM KH2PO4, pH 2.7, 30% MeCN) and separated by salt gradient consisting of 5 min at 100% solvent A, a 15 min gradient to 15% solvent B (solvent A + 0.35 M KCl), 1 min gradient to 100% B, 15 min at 100% B, and a 5 min gradient to 100% A. Fractions of 0.7 mL were collected throughout the run and combined into 8 fractions of equal peptide content based on their absorbance at 220 nm. Combined fractions were lyophilized prior to TiO2 enrichment.

TiO2 Enrichment of Phosphopeptides

Phosphopeptides were enriched using TiO2 in batch mode in the presence of 5% TFA and 1 M glycolic acid.13 Briefly, lyophilized peptides were resuspended in 80% MeCN, 5% TFA, 1 M glycolic acid prior to incubation for 15 min with TiO2 beads (GL Sciences) that were prewashed in the same solution. Beads were recovered by centrifugation at 700 × g for 1 min, washed three times in 80% MeCN, 5% TFA and eluted twice in 0.6% NH4 (aq). The combined eluents were lyophilized and stored at −20°C prior to analysis.

Mass Spectrometry Data Acquisition

Liquid chromatography tandem mass spectrometry was performed by the FingerPrints Proteomic Facility at the University of Dundee. Liquid chromatography was performed on a fully automated Ultimate U3000 Nano LC System (Dionex) fitted with a 1 × 5 mm PepMap C18 trap column and a 75 μm × 15 cm reverse phase PepMap C18 nanocolumn (LC Packings, Dionex). Samples were loaded in 0.1% formic acid (buffer A) and separated using a binary gradient consisting of buffer A (0.1% formic acid) and buffer B (90% MeCN, 0.08% formic acid). Peptides were eluted with a linear gradient from 5 to 40% buffer B over 65 min. The HPLC system was coupled to an LTQ Orbitrap Velos Pro mass spectrometer (Thermo Scientific) equipped with a Proxeon nanospray ion source. For phosphoproteomic analysis, the mass spectrometer was operated in data dependent mode with each MS scan triggering 15 MS2 acquisitions of the 15 most intense ions using multistage activation on the neutral loss of 98 and 145 m/z. Phosphopeptide data was processed using MaxQuant15 version 1.3.0.5 which incorporates the Andromeda search engine.16 Proteins were identified by searching a protein sequence database containing T. brucei brucei 927 annotated proteins (Version 4.0, downloaded from TriTrypDB,17 http://www.tritrypdb.org/) supplemented with the VSG221 sequence and frequently observed contaminants (porcine trypsin, bovine serum albumins and mammalian keratins) that contains a total of 10,081 protein sequences. Search parameters specified an MS tolerance of 6 ppm, an MS/MS tolerance at 0.5 Da and full trypsin specificity, allowing for up to two missed cleavages. Carbamidomethylation of cysteine was set as a fixed modification and oxidation of methionines, N-terminal protein acetylation and N-pyroglutamate were allowed as variable modifications. Phosphoproteomic analysis included phosphorylation of serine, threonine and tyrosine as additional variable modifications. Peptides were required to be at least 7 amino acids in length and a MaxQuant score >S, with false discovery rates (FDRs) of 0.01 calculated at the levels of peptides, proteins and modification sites based on the number of hits against the reversed sequence database. SILAC ratios were calculated using only peptides that could be uniquely mapped to a given protein group, and required a minimum of two SILAC pairs. To account for any errors in the counting of the number of cells mixed, the distribution of SILAC ratios was normalized within MaxQuant at the peptide level so that the median of log2 ratios is zero, as described by Cox et al.15

Prior to statistical analysis, the outputs from MaxQuant were filtered to remove known contaminants and reverse sequences, and phosphorylation sites with a MaxQuant localization probability <0.95 were discarded. This localization probability cut off was chosen following the manual inspection of the tandem mass spectra showing the best localization evidence for phosphorylation sites occurring on protein kinases.

Quantitation of the changes in abundance are derived from the two SILAC labeled experiments with a 1:1 mix of procyclic form cells grown in heavy-SILAC media and bloodstream cells grown in light-SILAC media, and the label swap experiment with the SILAC labeling reversed. SILAC ratios for phosphorylation sites were calculated using only data from the phosphoproteomic experiments, and SILAC ratios for proteins were calculated using only data from the nonenriched proteomic experiments. When SILAC ratios were reported in both label swap experiments, replicate trimming was applied by calculating the mean and standard deviation σ of the distribution of the log2 (H/L × L/H), and discarding data >2σ from the mean (Figure 3).18 Where SILAC ratios were reported in only one experiment, they were discarded if the percentage variation in the calculated SILAC ratio (calculated by MaxQuant) was >100%

The Mascot Delta Score was calculated by determining the difference between the best and second best Mascot ion scores for alternative phosphorylation site localizations on an otherwise identical peptide sequence,19 and appended to the MaxQuant outputs. Mascot distiller was used to peak pick mass spectra and read the centroid data from raw tandem mass spectra and submit searches to Mascot (Server version 2.2) using the same sequence database, enzyme specificity, modifications and mass tolerance as described above.

Data was visualized using Perseus 1.3.0.4 (www.perseus-framework.org) and further information on the identified proteins was obtained from TriTrypDB (http://www.tritrypdb.org).17 Gene ontology (GO) term enrichment was carried out using model based gene set analysis or term for term enrichment in Ontologizer20 using a T. brucei specific GO set containing 16765 terms for 4340 genes obtained from GeneDB (http://www.genedb.org).21 To make our data accessible to the scientific community, we uploaded our study to TriTrypDB and deposited the Thermo RAW files and search engine output into ProteomeXchange (http://www.proteomexchange.org) accession number PXD000049, enabling researchers to access the data presented here.

RESULTS

SILAC Labeling of Bloodstream Form Trypanosoma brucei

To enable SILAC labeling of bloodstream form T. brucei cells, a modified medium was created based on HM111-T (HM19-T
lacking serum plus), where L-arginine and L-lysine could be replaced by stable heavy isotope forms (HMI11-SILAC). As bloodstream form cells are usually only cultured to \(<2 \times 10^6\) cells/ml, a 10-fold lower density than procyclic cells, we first set out to establish labeling conditions that would reduce the amount of label required and maximize the number of cells obtained. We found that the growth of bloodstream form \(T. brucei\) cells in HMI11-SILAC containing only 30% of the normal concentration of L-arginine and L-lysine was equivalent to cell growth in the HMI9-T medium generally used for bloodstream form cell culture (Figure 1A). As expected, the HMI11-SILAC medium failed to support cell growth in the absence of L-arginine and L-lysine whereas cells were readily able to grow to \(5 \times 10^6\) cells/ml in the log-linear phase in the presence of L-arginine and L-lysine (Figure 1B). Furthermore, the gross morphology of the cells was unaffected after ten days in culture in HMI11-SILAC + 30% L-arginine and L-lysine, as judged by light microscopy (Figure 1C) and measurement of cell dimensions by light scattering (Supplemental Table 1, Supporting Information). Importantly, cells cultured in HMI11-SILAC + 30% L-arginine and L-lysine for >14 days maintained infectivity in mice (Supplemental Figure 1, Supporting Information). Taking all the above results into consideration, we concluded that culture of bloodstream form \(T. brucei\) in HMI11-SILAC + 30% L-arginine and L-lysine is equivalent to culture in HMI9-T medium.

To experimentally assess the efficiency of heavy isotope incorporation in HMI11-SILAC, bloodstream form cells were grown in HMI11-SILAC + \(\text{R}_6\text{K}_4\) for 11–12 cell divisions and extracted proteins subjected to analysis by LC–MS/MS. If heavy isotope incorporation occurred only by dilution (neglecting protein turnover), then 11–12 cell divisions should produce >99.9% incorporation, but in reality will be limited by the isotopic purity of the labeled amino acids. The heavy isotope incorporation at steady state was estimated to be 98.5 \(\pm\) 1.5% by comparing the relative abundance of the major isotopic peak of the heavy (arginine-\(^{13}\)C\(_6\)/lysine-\(^2\)H\(_4\)) and light forms of ten peptides chosen at random. No significant incorporation of proline-\(^{13}\)C\(_5\) (by conversion of arginine-\(^{13}\)C\(_6\)) was observed (Supplemental Figure 2, Supporting Information), as noted previously for the procyclic form.3

**Comparative Proteomic Analysis**

A prerequisite for comparative quantitative phosphoproteomic analyses is access to a relevant comparative quantitative proteomic data with which to distinguish the different types of quantitative phosphorylation differences (i.e., those that simply track changes in protein levels, those that do not correlate with changes in protein level and those that are unique to one or other lifecycle stage). Our previous comparative SILAC proteomic analysis of bloodstream grown in light medium and procyclic form \(T. brucei\) grown in heavy medium gave us ratios for 3553 proteins, and demonstrated that large changes in specific protein abundance occur between the lifecycle stages.3 To improve the quality of this comparative quantitative proteome data set, we exploited the development of SILAC labeling of

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**Figure 1.** Growth of bloodstream form \(T. brucei\) in SILAC labeling media. Cells were grown in conventional HMI9-T medium or in the HMI11-SILAC medium developed in this paper. (A) Cumulative growth over 10 days (with subculturing every 2–3 days). (B) Growth to stationary phase. (C) Light microscopy of cells after 10 days in culture.
cells grown in light-SILAC medium. (ii) A 1:1 mix of procyclic form cells grown in heavy-SILAC medium and bloodstream form cells grown in light-SILAC medium. (iii) A “label swap” experiment consisting of a 1:1 mix of bloodstream form cells grown in light-SILAC medium and procyclic cells grown in heavy-SILAC medium. (iv) Bloodstream form cells grown in light-SILAC medium. To maximize coverage of membrane and structural proteins, total protein extracts were prepared with 4% SDS using the filter-aided sample preparation (FASP) procedure, resulting in near-complete solubilization of the parasites. After denaturation and reductive alkylation the proteins were digested with trypsin in solution. Resulting phosphopeptides were enriched and fractionated into eight samples by SCX chromatography prior to further enrichment using TiO2 chromatography in batch mode. For each phosphoproteomic experiment, the eight fractions were analyzed by liquid chromatography tandem mass spectrometry in technical triplicates on a LTQ Orbitrap Velos using multistage acquisition.

### Overview of Data

The combined data were from a total of 138 LC−MS/MS runs, with 96 runs for samples enriched for phosphopeptides and 88 for samples with SILAC labeling. Altogether 589101 MS/MS spectra were identified, corresponding to 58087 nonredundant peptide sequences belonging to 5478 protein groups with a false discovery rate of 1% (Supplementary Tables S2 and S3, Supporting Information). A total of 10096 phosphorylation sites were identified on 2551 proteins (Supplementary Tables S4 and S5, Supporting Information), of which SILAC ratios could be determined for 8275 phosphorylation sites on 2233 protein groups, with a Pearson correlation of 0.78 between the label-swap experiments. A total of 8120 phosphorylation sites on 2137 protein groups were also identified using Mascot software, of which 6196 phosphorylation sites on 1933 protein groups had a Mascot delta score ≥10, reported to give a 1% false localization rate (FLR) for Orbitrap-MSA data. Of the 10096 phosphorylation sites identified by MaxQuant, 9314 (92%) have not previously been reported. The data presented here therefore represents a major advance in the depth of the trypanosome phosphoproteome.

The distribution of phosphorylation between residues was found to be 8087 phosphoserine (pS) sites (80%), 1,720 phosphothreonine (pT) sites (17%), and 289 phosphotyrosine (pY) sites (3%), in reasonable agreement with the *T. brucei* bloodstream form cytosolic phosphoproteome observations of 75% pS, 21.5% pT, 3.5% pY. Of the 10096 phosphorylation sites observed, 6493 occurred on 1491 proteins annotated as hypothetical or hypothetical conserved, reflecting the high proportion of genes of unknown function in the trypanosome genome. Gene Ontology terms are associated with only 1310 out of the 2552 phosphoproteins we observed, limiting the power of GO term enrichment analysis. Performing model based gene set analysis, a method designed to give a high-level summarized view of the data, identified only three terms: ciliary or flagella motility (GO:001539, PEP 1), protein tyrosine kinase activity (GO:004713, PEP 0.7), and regulation of gene expression (GO:0010468, PEP 0.5). This is an intriguing outcome as the *T. brucei* genome lacks any identifiable dedicated tyrosine kinases, yet has the associated GO terms, illustrating the pitfalls of such analyses. Performing term-for-term enrichment generated 45 significantly enriched GO terms (p < 0.01, Figure 4) confirming the predominance of terms associated with flagella motility, protein kinase activity and regulation of gene expression, and revealing additional terms for RNA, zinc and calmodulin binding (GO:0005488, GO:0008270, GO:0005516), and peptidase activity (GO:0004181, GO:0004197).

Greater biological insight from the data is likely to come from the community inspecting the phosphorylation patterns, and their changes between lifecycle stages, of proteins of specific interest to them, a few examples of which are provided below.

![Figure 2](https://example.com/figure2.png)

**Figure 2.** Phosphoproteomic workflow. Differentially labeled (heavy and light) bloodstream and procyclic form trypanosomes were lysed in SDS, and total proteins S-alkylated and digested with trypsin using the FASP method. Phosphopeptides were separated into eight fractions by SCX HPLC, enriched with TiO2, and analyzed by LC−MS/MS using MSA.
Comparison of the Phosphoproteomes of Bloodstream and Procyclic Form *T. brucei*

SILAC labeling allows the fold-changes that occur between the bloodstream and procyclic form of *T. brucei* (log2 Pcf/Bsf) to be quantified at both the protein and phosphorylation site level. As the phosphoproteomic workflow includes selective enrichment for phosphorylated peptides, this data cannot be used in the calculation of protein SILAC ratios, which are instead derived from separate proteomic experiments. We were able to determine fold-changes for a total of 8275 phosphorylation sites on 2233 protein groups, with 5768 (70%) of the phosphorylation sites on 1385 protein groups having ratios at both the protein- and site-level. In general, the changes in phosphorylation site occupancy between the lifecycle stages are greater than the changes in protein levels; for example, 318 (5.5%) of the phosphorylation sites change ≥10-fold (log2 ≥3.32) between lifecycle stages, while only 33 (0.87%) of protein groups change ≥10-fold. Indeed, plotting the fold-changes in protein ratio against the fold-changes in phosphorylation site ratio (Figure 5A) reveals that the observed changes are relatively weakly correlated (Pearson correlation 0.49). However, there are numerous examples where: (i) The change in phosphorylation correlates with the change in protein abundance, suggesting that phosphorylation stoichiometry is maintained between lifecycle stages in these cases. (ii) The change in phosphorylation correlates negatively with the change in protein abundance. (iii) Protein abundance is maintained but phosphorylation stoichiometry changes significantly (both up and down). (iv) Different phosphorylation sites on the same protein show a wide range of fold-changes (Figure 5B and Figure 6). These data demonstrate that changes in protein phosphorylation status between the lifecycle stages of *T. brucei*...
add significant complexity to the differential expression of the proteome. Some individual cases are discussed in more detail below.

**Manual validation of protein kinase phosphorylation sites**

In order to judge the quality of the phosphorylation site assignments, and to empirically assess the false localization rate, a subset of the automated phosphorylation site assignments were manually inspected (Figure 6 and Supporting Information). The subset chosen was the *T. brucei* kinome consisting of 176 protein kinases as they are expected to occur at relatively low abundance and to show physiologically relevant phosphorylation. Using a MaxQuant localization score of $\geq 0.75$ as an initial cutoff, we identified 641 specific phosphorylation sites on 127 protein kinases (for comparison a cutoff of $\geq 0.95$ gave 493 sites on 124 kinases). Of these 641 automatically annotated sites on 127 kinases, 549 sites on 122 kinases were validated by manual inspection (Supplemental Table S6, Supporting Information). Overall, the manual validation correlated well with the sites chosen using the MaxQuant localization score and the Mascot delta score (MDS) methods. However, we noted that only 42% of sites in the MaxQuant localization score range $0.75 - 0.95$ were manually validated, whereas over 94% of the sites in the MaxQuant localization score $>0.95$ were manually validated, increasing to over 98% at a localization score of $>0.99$. Further, at an MDS of 10, reported to give a global 1% false localization rate for Orbitrap Velos – MSA data, we were able to manually confirm 98% of the automatically annotated sites. However, it is worth noting that 81% of sites with MDS < 10 and 66% of sites identified by MaxQuant but not by Mascot were manually validated. Based on this manual survey of a significant number of phosphorylation site assignments, we have chosen to report all automatically annotated phosphorylation sites with MaxQuant localization scores $>0.95$ (with an estimated false localization rate of <6%).

Interestingly, the variation in the phosphorylation of the kinome between the two lifecycles stages is lower than the general data set, with only 15 phosphorylation sites showing a 10-fold or greater change. The protein kinase A catalytic subunit isoform 1 (Tb09.211.2410) showed strongly up-regulated phosphorylation in the bloodstream form parasite at three sites ($\log_2$ range $-5.0$ to $-4.3$), although the assignment of increased phosphorylation in the putative activation loop at T179 ($\log_2 -2.6$) is slightly ambiguous as an identical (phospho)peptide is shared by protein kinase A catalytic subunit isoform 2 (Tb09.211.2360). Similarly, the protein kinase A regulatory subunit (Tb11.02.2210) has 5 sites which are also up-regulated in the bloodstream form, though to a lesser extent ($\log_2$ range $-4.4$ to $-0.8$). Protein kinases with phosphorylation up-regulated in the procyclic form include a NEK kinase (Tb11.01.6650) with two sites with a $\log_2$ range 5.6 to 4.7, and two CLK dual specificity kinases Tb927.2.4200 (5 sites, $\log_2$ range 3.3 to 0.6) and Tb927.10.15020 (4 sites, $\log_2$ range 3.2 to 2.4). Two “Other” group protein kinases have phosphorylation sites that are up-regulated in the procyclic and bloodstream form, respectively, (Tb927.7.3210, 5 sites, range $\log_2$ $-3.7$ to 2.8; Tb927.7.3220, 5 sites, range $\log_2$ $-3.2$ to 1.6).

Of the 15 MAPKs present in the *T. brucei* kinome, 9 (ECK1, KFR1, MOK, RCK, MAPK2, MAPK9, MAPK10, MAPK12, and an unnamed MAPK) are shown by the data presented here to have dual phosphorylation in the TxY motif required for activation. In addition, multiple phosphorylation sites are observed on three potential upstream STE7 MAP2K and nine STE11 MAP3K protein kinases, suggesting the presence of active MAPK signaling cascades. The observation of differential phosphorylation on certain MAPKs, MAP2Ks and MAP3Ks between the two life-cycle stages suggests that MAPK signaling may play distinct roles in each lifecycle stage (Figure 7).

**Phosphorylation Sites Up-regulated in Bloodstream Form *T. brucei***

A total of 203 phosphorylation sites on 119 protein groups were up-regulated $\geq 10$-fold ($\log_2 \leq -3.32$) in bloodstream form *T. brucei*, with a distribution of these up-regulated phosphorylation sites...
sites of 74% pS, 15% pT, 11% pY, showing a significant enrichment in pY compared to the total observations. An additional 429 sites with no SILAC ratio (on 319 protein groups) were observed exclusively in the bloodstream form, with the distribution of 64% pS, 27.5% pT, 8.5% pY showing a similar trend. These data suggest that tyrosine phosphorylation is significantly

Figure 6. Examples of the assignments of phosphorylation site location from tandem mass spectral data. (A) STE11 family protein kinase (Tb11.46.003) with pS357 and pS358 with MaxQuant localization score of 1.0 and Mascot delta score of 57.8. (B) CAMK family protein kinase Tb11.01.2290 with pS5 with MaxQuant localization score of 1.0 and Mascot delta score of 27.1, and pS8 with MaxQuant localization of 1.0 but which was not identified by Mascot. Data acquired on a LTQ Orbitrap Velos using MSA, * designates loss of phosphate (−98 Da), ** loss of two phosphates (−196 Da).
over represented in sites differentially regulated between the lifecycle stages.

In total, 439 proteins have 631 individual phosphorylation sites that are either ≥10-fold up-regulated in the bloodstream form or are bloodstream form specific (Supplemental Table S7, Supporting Information). This group contains a large number of proteins annotated as hypothetical or hypothetical conserved, including 13 of the 15 proteins with >40-fold change (Log2 < −5.32) at the site level. GO term enrichment of the bloodstream form up-regulated group (of the 191 annotated proteins out this total 439) produced a very similar set of enriched terms to the global and bloodstream form up-regulated phosphoproteome (flagella, protein kinase, and gene expression), with the addition of terms for glycolysis and gene expression), with the addition of terms for glycolysis and energy metabolism, although this may be due, at least in part, to large changes at the protein level in some of these proteins. Nevertheless, researchers interested in the changes in parasite metabolism between the procyclic and bloodstream form parasites are likely to find clues to physiologically relevant phosphorylation changes in the data provided. Significantly, there are examples where protein up-regulation in bloodstream form parasites may be further “amplified” by the coordinate up-regulation of phosphorylation. Notable examples of this include the RNA recognition motif protein RBP10 (Tb927.8.2780) that has been reported to promote a bloodstream form mRNA expression pattern, which has 4 phosphorylation sites changing by an average of log2 < −4.8 (from log2 < −5.4 to −4.2), potentially amplifying a change in protein abundance of log2 < −3.3, and the receptor-type adenylate cyclase GRESAG4 (Tb927.8.7940) which has 6 phosphorylation sites changing by an average of log2 < −3.9 (from log2 < −4.7 to −3.1), potentially amplifying a change in protein abundance of log2 < −2.4.

**Phosphorylation Sites Up-regulated in Procyclic Form T. brucei**

A total of 115 phosphorylation sites on 83 protein groups were ≥10-fold up-regulated (log2 ≥ 3.32) in procyclic form T. brucei with a residue distribution of 70% pS, 22% pT, 9% pY, and an additional 499 sites with no SILAC ratio on 376 protein groups were observed exclusively in the procyclic form with a residue distribution of 77% pS, 15.5% pT, 7.5% pY. The significant enrichment in pY compared to the total observations mirrors that seen in the bloodstream form up-regulated phosphorylation sites, reinforcing the importance of changes in the phosphorylation of tyrosine between the lifecycle stages.

In total, 459 proteins have 614 individual phosphorylation sites that are either ≥10-fold up-regulated in the procyclic form or are procyclic form specific (Supplemental Table S8, Supporting Information). This procyclic up-regulated group contains a large number of proteins annotated as hypothetical or hypothetical conserved, as observed for the bloodstream form up-regulated group. GO term enrichment of the procyclic up-regulated group (of 207/460 proteins) again produced a very similar set of enriched terms to the global and bloodstream form up-regulated phosphoproteome (flagella, protein kinase, and gene expression), with the addition of new terms. A total of 9 proteins annotated as transporters have >10-fold change (log2 > 3.32) at the site level, potentially amplifying the corresponding changes in protein abundance. Interesting specific examples include: The PAD2 carboxylic acid transporter (Tb927.7.5940), associated with slender to stumpy bloodstream form transformation in pleomorphic cell lines, where 3 sites increase in phosphorylation with an average log2 of 3.3 (range log2 1.5 to 6.2) in addition to a protein level increase of log2 2.9. The ubiquitin carboxy-terminal hydrolase (Tb927.10.2210), which has 10 phosphorylation sites with an average log2 of 3.6 (range log2 1.6 to 5.6), while the protein has almost no change in level (log2 < −0.41). The C-14 sterol reductase (Tb11.01.7170), involved in the biosynthesis of ergosterol and its isomers that are the principal sterols in the procyclic parasites, has 2 phosphorylation sites with an average log2 of 3.8 (range log2 3.7 to 3.9), potentially amplifying the protein fold change of log2 2.2.

**Phosphoproteins with Differentially Regulated Sites**

A significant number of proteins that are multiply phosphorylated exhibit phosphorylation sites that are differentially regulated within the protein between the lifecycle stages (Figure 8). There are

**Figure 7. Differential phosphorylation of MAPK signaling components.** Heatmap showing the change in abundance between the procyclic to bloodstream (Log2) at the protein and phosphorylation site level for the MAPK, STE7 and STE11 kinase quantified in this study. Red box: TxY motif; Gray shading, not observed. The Heatmap is generated with GENEE (http://www.broadinstitute.org/cancer/software/GENE-E/).

**Figure 8. Distribution of intraprotein differentially regulated phosphorylation sites.** Only proteins with more than one phosphorylation site are shown.
enrichment analysis (of 113/234 proteins) produced significantly enriched terms (p < 0.01) including flagellum, gene regulation, and protein phosphatase activity, but that did not include protein kinase activity. Three protein phosphatases are found in the intraprotein differentially regulated group: (i) A kinetoplastid-specific protein phosphatase (Tb11.01.4320) with a differential range log₂ of 6.6 (10 sites, range log₂ −4.8 to 1.8). (ii) An atypical dual specificity phosphatase (Tb11.01.5870) with a differential range log₂ of 5.6 (3 sites, range log₂ −3.8 to 1.7). (iii) The TFIIF-Stimulated CTD phosphatase (Tb927.10.11450) with a differential range log₂ of 4.1 (4 sites, range log₂ −2.1 to 2.0).

## DISCUSSION

We have previously developed SILAC labeling in procyclic form *T. brucei*, and here we report the development of SILAC labeling in the clinically relevant bloodstream form and demonstrate that growth, morphology and infectivity are unchanged. Both lifecycle stages are rapidly dividing, enabling efficient stable isotope incorporation >98%, and show no propensity to convert arginine to proline. Two recent proteomic analysis using SILAC labeling in either procyclic or bloodstream form also observed a similar lack of arginine to proline conversion.

The quantitation of changes in the global phosphoproteome between procyclic and bloodstream form lifecycle stages of *Trypanosoma brucei*, described here, reveals that differential phosphorylation is widespread, is not strongly correlated to change in protein abundance and adds significant complexity to the differential expression of the proteome. The observation of intraprotein differential phosphorylation suggests that phosphorylation may play distinct functional roles in each lifecycle stage, although how this effect is mediated is at present unclear and, because of the lack of comparative phosphorylation data to date, there are few existing hypotheses to examine. However, one process where protein phosphorylation is suggested to play a central role is the differentiation of the parasite from bloodstream to procyclic form in a coordinated, programmed event. While we have not studied the actual process of differentiation in this study, we have examined the phosphoproteomes of the initial and final states of the process, and observe phosphorylation changes in many of the proteins involved in differentiation. For example, TOR4 (Tb927.1.1930) has 10 sites with log₂ changes in the range 0.4−2.7, MAPKS (Tb927.6.4220) has a protein change log₂ of 0.5 and pT207 site change log₂ of 0.8, ZFK (Tb11.01.1030) has a pSS20 log₂ of −1.3; NOPP44/46 (Tb927.8.760) has a protein log₂ of 1.7 and pY5 observed uniquely in procyclic parasites; PAD2 (Tb927.7.5940) has a protein log₂ of 2.9 and 3 sites with log₂ in the range 1.5 to 6.2. The absence of detected phosphorylation on PIP3 (Tb09.160.4480, protein log₂ of 0.9), and PTP1 (Tb927.10.6690, not observed in our data set) may be due to up-regulation or transient phosphorylation during differentiation, such as in the division arrested stumpy form that was not studied here. Alternatively, the cocktail of phosphatase inhibitors used in this study may have limited effect against the activity of one or more trypanosome phosphatase. Clearly, further detailed studies of the process are required before firm conclusions can be drawn on the precise rules of protein phosphorylation in trypanosome differentiation.

In this study we have used culture adapted Lister 427 bloodstream form (MITat 1.2) and procyclic form (29.13.6) cell lines that have been made amenable to genetic manipulation by Wirtz et al. by the insertion of exogenous genes coding for the T7 RNA polymerase and tetracycline repressor protein using either one (bloodstream form) or two (procyclic form) selectable drug resistance markers. The two cell lines used are independent clones that are not genetically identical, and this may lead to some differences in protein expression and phosphorylation status, although such differences are likely to be minor in comparison to the developmentally regulated changes between the lifecycle stages. In addition, the culture adapted Lister 427 strains are neither genetically nor morphologically identical to field isolates, and it is not possible to account for the impact this will have on protein expression and phosphorylation status.

A significant proportion of the *T. brucei* kinome of 176 protein kinases are phosphorylated, with individual examples of phosphorylation sites up-regulated in either procyclic or bloodstream form, and examples of intraprotein differential phosphorylation. The abundance of active MAP kinases, as judged by the presence of dual TxxY motif phosphorylation, suggests that MAPK pathways are functional in both lifecycle stages of *T. brucei*, although how or whether the pathways are regulated remains unclear. Efforts to determine the detailed biological role of *T. brucei* protein kinases are ongoing and knock-down by RNA interference has provided evidence of the essential nature of a significant number of protein kinases. We have recently used chemical proteomic profiling to establish that *T. brucei* protein kinases are sensitive to typical kinase inhibitors with nanomolar potency, demonstrating the potential for the development of species-specific inhibitors. Further analysis of protein kinase mediated phosphorylation cascades by stimulus−response studies, using the data published here to anchor them, may reveal the pinch-points that are most suitable for therapeutic intervention.

Often, eukaryotic phosphorylation signaling cascades terminate in the modulation of transcription factors to mediate changes in gene expression. However, classical transcription factors are missing in *T. brucei*, where the processing and stability of polycistronically transcribed mRNA precursors appears to regulate gene expression. One potential mechanism to modulate gene expression in *T. brucei* is the use of RNA binding zinc finger proteins to influence mRNA processing and stability, such as such as ZFP3 (Tb927.3.720) and RBP10 (Tb927.8.2780). A significant number of ZFPs (31 out of 49) and RBPs (15 out of 47) are present in the phosphoproteomic data reported here, with a subset of these (11 and 5, respectively) showing lifecycle specific regulation of phosphorylation status (≥5-fold change). Since the dynamic phosphorylation/dephosphorylation of these proteins has the potential to modulate gene expression, they are reasonable candidates to be the ultimate effector molecules of at least some of the trypanosome signaling cascades.

Despite the lack of identifiable dedicated tyrosine kinases in *T. brucei*, 234 phosphotyrosine residues were identified, and tyrosine phosphorylation was 3−4 fold over-represented in the phosphorylation sites that change between the two lifecycle stages. Tyrosine phosphorylation is proposed to arise from putative dual-specificity protein kinases such as DYRK, MEK and CLK, but there has been no experimental verification of the dual specificities of the trypanosome kinases to date.

GO term enrichment analysis has revealed that the flagellum is consistently over-represented among the phosphorylated proteins. Flagella can play both motility and sensory roles, and while it is clear that the trypanosome flagellum is highly motile, its potential sensory role is less well-defined. The dynamic phosphorylation of dynein motors is proposed to control flagella motility in other organisms, and in trypanosomes flagella motility is dispensable in the procyclic form but essential for cytokinesis in the bloodstream form. Interestingly, a number of flagella
proteins show extensive intraprotein differential phosphorylation, including: (i) dynein heavy chain (Tb927.7.920) with 7 sites with log2 changes in the range −2.8 to 3.2 and protein log2 changes of −0.5; (ii) a paraflagella rod component (Tb927.8.1550) with 14 sites with log2 changes in the range −3.1 to 2.0 and a protein log2 change of −0.4; and (iii) an outer dynein docking complex (Tb11.01.7750) with 3 sites with log2 changes in the range −3.6 to 1.8 and a protein log2 change of 0.6.

■ CONCLUSIONS

In summary, we have presented a global quantitative phosphoproteomic analysis of the bloodstream and procyclic form of *Trypanosoma brucei*, which represents both the largest set of phosphorylation sites and the first report of genome-wide quantitative phosphorylation data for any kinetoplastid. The establishment of SILAC labeling in both culture-amebic lifecycle stages of the parasite provides the community with the tools required to examine signaling events at the molecular level. This data set reveals the breadth of protein expression and phosphorylation in *T. brucei*, and provides a resource for the community to begin to examine the effect of phosphorylation upon biological processes. The present study of rapidly dividing nonsynchronized cells provides a static, averaged snapshot of the basal level of protein phosphorylation, and valuable information will be gained from future studies of the dynamics of stimulus-response events such as differentiation.

■ ASSOCIATED CONTENT

◆ Supporting Information

Assignment of MS2 spectra for protein kinase phosphorylation sites, as pdfs contained in a zipped file. Supplementary Figures S1. Rodent infectivity of bloodstream form *Trypanosoma brucei* cultured in HM9-T and HI11-SILAC media. Figure S2. Efficiency of isotope incorporation in bloodstream form *T. brucei*. Figure S3. Gene Ontology term enrichment for all phosphoproteins. Table S1. Cell dimensions. Table S2. Protein groups including SILAC ratios. Table S3. Peptide including SILAC ratios. Table S4. Phosphorylation sites including SILAC ratios. Table S5. Phosphopeptides including SILAC ratios. Table S6. Manual validation of protein kinase phosphorylation sites. Table S7. Phosphorylation sites up-regulated in the bloodstream form. Table S8. Phosphorylation sites up-regulated in the Procyclic form. Table S9. Phosphoproteins with phosphorylation sites differentially regulated between lifecycle stages. This material is available free of charge via the Internet at http://pubs.acs.org.

■ AUTHOR INFORMATION

Corresponding Author

*E-mail: m.a.j.ferguson@dundee.ac.uk. Tel.: 44-1382-384219. Fax 44-1382-245764.

Notes

The authors declare no competing financial interest.

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■ABBREVIATIONS

FASP, filter aided sample preparation; GO, gene ontology; SCX, strong cation exchange; SILAC, stable isotope labeling by amino acids in cell culture; T. brucei, *Trypanosoma brucei*

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