A Step-By-Step Guide for Reducing Time-Series Photometry with the HiPERCAM Pipeline for the Boston University White Dwarf Group

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1 Conventions for this document

1.1 Fonts

Text written in typewriter font and enclosed in thistle-colored boxes is generally meant to be copy and pasted into the command line, with individual commands being separated from one line to the next. For example:

```
mkdir test_directory cd test_testdirectory
```

1.2 Disclaimer on Hardware and Software

For posterity, I developed this document and the instructions herein working from my MacBook (MacOS=v15) whose Terminal operates using Zsh, not bash. While the necessary adjustments from Zsh to bash should be trivial, it is unclear how installation will be affected from MacOS to Windows, for example.

2 Installation and Configuration

2.1 Installing hipercam

You must first install and configure the HiPERCAM pipeline on your machine before you can crank out any light curves.

This is best done by creating a new conda environment¹:

```
conda create -n hipercam python=3.9.19
```

You can then install the pipeline as directed on https://github.com/HiPERCAM/hipercam:

```
mkdir hipercam
cd hipercam
conda activate hipercam
git clone https://github.com/HiPERCAM/hipercam.git
pip install . --user
```

2.2 Installing pgplot

You must next install the dependencies for hipercam. These include pgplot. For Mac users, this is best achieved through homebrew: follow these instructions.

Next you must install Tom Marsh's Cython wrapper to pgplot: follow these instructions.

2.3 Installing phot21c

All still within your hipercamconda environment, you should next install <u>phot21c</u>, Zach Vanderbosch's light curve extraction tool designed to replace WQED:

 $^{^{1}}$ The selection of this Python version is arbitrary. I have not tested this on more recent version releases (v≥3.10). At the time of developing this pipeline, v3.9.19 was the most up-to-date version of Python 3.9.

```
pip install phot21c
```

You next need to configure phot21c to be compatible with our hipercam-reduced photometry: photconfig

If you happen to be lucky enough to be Joseph Guidry and reducing PTO+PRISM photometry on his machine, your inputted parameters would be:

```
author
                  = Joseph Guidry
image_list_name
                  = hcm.lis
pixloc_name
                  = aperture.ape
                  = output
photbase_name
stardat_location = /Users/astrojoe/Research/Variability/phot2lc/stars.dat
default_telescope = pto
default_source
                  = hcm
default_image
                  = None
                  = None
default_object
```

See https://phot2lc.readthedocs.io/en/latest/configuration.html for more information. stars.dat is simply a text file with three columns: the object's name, and then its RA and dec in HH MM SS.SS +/-DD MM SS.SS formatting.

Finally, for users of PTO+PRISM and LDT+LMI, see Appendix ?? for the necessary hard-code to pasted into to the teledat.py file within phot21c.

2.4 Install Remaining Dependencies

It is likely you will not have all the requisite Python packages installed to run phot2lc, and perhaps even hipercam. You likely are lacking—based on my personal experience—pandas:

```
And lmfit:

pip install lmfit

And possibly even PyQt5:

pip install PyQt5
```

Do check the requisite dependencies for <u>hipercam</u> and <u>phot21c</u> and compare against what you currently have installed (run conda list in Terminal) to be most prudent.

You probably also need to have XQuartzinstalled to be able to use PGPLOT.

The final Python package to install is <u>ccdproc</u>, which is used in the Python image reduction routine calibrate_science_images that I custom-built into hipercam:

```
conda install -c conda-forge ccdproc
```

2.5 Configuring hipercam

2.5.1 Updating your Zsh Path

After every hipercam installation², make sure "/usr/home/.local/bin" is in path so you can execute the hipercam command line commands. Again, to use my machine as for illustration, this check would look like:

```
echo $PATH
path+=(/Users/astrojoe/.local/bin) to append OR
path=(/Users/astrojoe/.local/bin $path) to prepend
export PATH
source ~/.zshrc
```

hipercam will inform you of the exact directory you need to append into your path, which will vary from user-to-user. See this stackoverflow post for more information on this process of updating your Z shell path. I've had issues with keeping this path amendment permanent, so you may need to re-append after a reboot.

2.5.2 Updates to hipercam Hard Code

To be able to run this pipeline as instructed, I had to make various amendments to the hipercam source code. From this GitHub repo, download the following edited files I prepared to replace the originals within in your hipercam directory:

- fits2hcm.py
- setaper.py
- calibrate_science_images.py this is a novel routine I wrote
- setup.py

Each file should be uploaded to '~/hipercam/hipercam/scripts/', except setup.py, which belongs in the root directory, '~/hipercam/'.

You must re-install hipercam so these edits to the source will take effect (again, still with your hipercam conda environment activated AND within your hipercam directory):

```
pip install . --user
```

You also need to replace a few phot21c files. These should be uploaded in, using my machine again as an example: '/Users/astrojoe/Python/anaconda3/envs/hipercam/lib/python3.9/site-packages/phot2lc/'. The files to download are:

- photfunc.py
- teledat.py

3 Calibrate your Science Images

You now have all the necessary software installed and configured on your machine. It's now time to calibrate your raw photometry. We achieve this by running: calibrate_science_images --i <instrument>

²You must re-run pip install . --user within your hipercam directory after making any edits to the source code for the updates to take effect.

The instrument choices are ProEM, PRISM, and LMI (or another instrument that hipercam is compatible with).

This routine works form the assumption that you have organized your nightly data in a directory structure such as the following:

- bias
- dark (if applicable for your instrument!)
- dome_flat or sky_flat
- WDJHHMM+/-DDMM it's not necessary to name your folder after the white dwarf's coordinates, but you science images should be in some unique folder separate from your calibration images. You should also have a unique folder for every target observed.

These all would live in a directory named after the UTC date of the night you observed: 'YYYYM-MDD/', which again is not essential, but is a wise convention to adopt.

This structure is necessary because you must run calibrate_science_images (and all the remaining hipercam routines) within the folder of the target whose photometry you are attempting to reduce. Once commanded, calibrate_science_images will prompt you to edit the header information of your science images and check the timestamps. It will then produce master calibration images (if they have not yet been made from reducing another object of this night), and then bias subtract, dark subtract (if applicable), and flat field your raw science images, finally saving them as new copies, now with 'c.fits' extensions.

calibrate_science_images also writes out two files: aperture.ape and reduce.red, which will be used below, along with creating a new directory for you .hcm files that you will soon write out. Before reducing and extracting your photometry, you should be choose how you wish to extract your photometry. By default the file is created to do PSF photometry using variable apertures in the "optimal" mode. You can instead do fixed aperture photometry, the "normal" mode. You can change between these modes in a text editor. A meticulous astronomer would reduce their photometry using both methods, along with tinkering with other settings, to achieve the highest fidelity photoemtry. Refer to Tom Marsh's notes for more information:

https://cygnus.astro.warwick.ac.uk/phsaap/hipercam/docs/html/photometry.html.

3.1 Known Bugs

When you execute this calibrate_science_images, an ImportError will be prompted upon completion. **Disregard this. The routine successfully completed.** I don't know how to eliminate this. It stems from me forcing this into the source code.

A second bug that I hope to resolve is when you took images in multiple filters, the multiflats function can get hung up. You might be prompted to enter the path to your flats directory. Enter it. It will probably fail. Don't fret. Re-run the routine and it should complete successfully.

4 Reduce your Photometry with hipercam

With our calibrated science images, we can finally perform our photometry and extract light curves. But first, we must convert our 'c.fits' images into file formats that hipercam can read. We do this by running fits2hcm:

```
fits2hcm hcm.lis [origin] True
```

Here, origin is either "PRISM", "ProEM", or "LMI".

UPDATE ADDITIONS TO FNAME, BNAME, ONAME AND PROEM AND LMI COMPATIBILITY

Next, we have to set the aperture positions of our target and comparison stars using setaper:

```
setaper hcm_files/[first image] aperture.ape 10 20 30 xlo xhi ylo yhi True p 1 99
```

In the pop-up window, mouse over your target and type the 'a' key. Repeat for each comparison star.

It can be prudent to trim the image to exclude the margins from consideration. For example, a 410×380 PRISM image could be trimmed to xlo=10, xhi=400, ylo=10, yhi=370. ProEM images should use the native dimensions: 0 256 0 256. If your target is too faint for the PSF to be identified, try reducing the range percentile range to 5 95.

Finally, run reduce to perform the photometry extraction:

```
NEED TO make COPIES OF REDUCE.RED for each instrument reduce source=hf hcm.lis False reduce.red output.log 0 True False
```

The final 'False' parameter can be changed to 'True' to prompt a window to pop up that projects each image and the apertures as you are performing the photometry from one image to the next. It is best practice to monitor this in case a aperture begins to wander away from a target or comparison star and begin to randomly walk around the images. This will only occur if the source is extraordinarily faint. You can guard against this by reducing the box size to search for Gaussian sources within when the variable aperture positions are being searched for: reduce the 'search_half_width' value in reduce.red. If some comparison stars prove to be problematic, either because they are so bright they go non-linear, or wander too near the edge of the chips, for example, re-run setaper and then re-run reduce.

5 Extract your Light Curve with phot21c and Post-Processing

Ta-da! You reduced your photometry. Now we need to extract our light curve, clean it up with some processing, barycentric-correct the timestamps, and write it out, all using the phot2lc suite.

To extract your light curve with phot21c simply call (still in that hipercam conda environment!):

phot21c [-t --telecode] [-o --object]

For PTO photometry, you may have to include the object name: WDJHHMM±DDMM, where HHMM and DDMM are the sexagesimal RA and dec coordinates in hours and minutes and degrees and minutes, respectively.

Relevant telecodes to BUWD observers:

- pto Perkins Telescope Observatory, set as default (§ 2.3)
- mcd McDonald Observatory 2.1-m Telescope
- 1dt Lowell Discovery Telescope

When using phot21c, consult Zach's documentation for all the commands and best practices: https://phot21c.readthedocs.io/en/latest/quickstart.html#basic-usage

You can save a quicklook plot of the light curve you extracted using the phot21c script quicklook: quicklook [-f --.lc filename] [-s --save plot] [-p --prewhiten periodogram]

Another useful post-processing script is weldlc. Say you had two runs within the same night of the same target, but had to change exposure times because of clouds. You can combine these into a single light curve file using weldlc, which will correctly order the timestamps from the barycentric corrections that phot2lc applies. You can weld together as many light curves as you wish, aggregating years-worth of photometry of a given star if you wish.

weldlc [-f --.lc filenames] [-o --output filename]