



# Lecture 8:

## Research posters

### Purpose and design

**Course 501**

Writing and Communication Skills



# AIMS

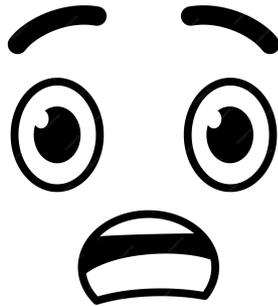
- Learn about the purpose of scientific posters.
- Understand the basic elements of posters.
- Review recommendations regarding poster design.
- Design your own poster based on the material of your selected research paper.

**IMPORTANT!**

# Scientific posters

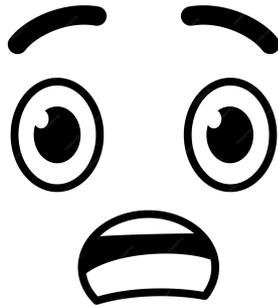
## Shocking truths

**Your poster is not**



a scientific paper  
a peer-reviewed material  
a medium for scientific  
debate  
a one time work  
a major scientific  
contribution  
a proof scientific quality  
a source of stress  
to convince others  
your *magnum opus*

**Your poster is**



to say I am scientifically alive  
to say I am still working  
to socialize  
to collaborate  
to get new ideas  
to modify your ideas  
to get help  
to get soft criticisms  
to enhance your science  
a frequent event  
a reason to leave the lab

**IMPORTANT!**

Scientific content  
Keep in mind



Your poster may not be as developed and complete  
as your future scientific paper.

Your poster can be as simple as a layout of future  
experiments.



Your poster does not need to follow the standard sections (Abstract, Introduction, M&M etc.).

You may be tempted to follow the layout of scientific papers due to its wide use in design.



The text content of your poster does not need to be in a paragraph format.



Use bullet points of concise short sentences rather than long and complicated paragraphs.



**Too much text** is not attractive, a sign of sophistication, or mirrors high-quality work.

**Too much text** is repulsive to the audience and a sign of desperation to fill the poster area.

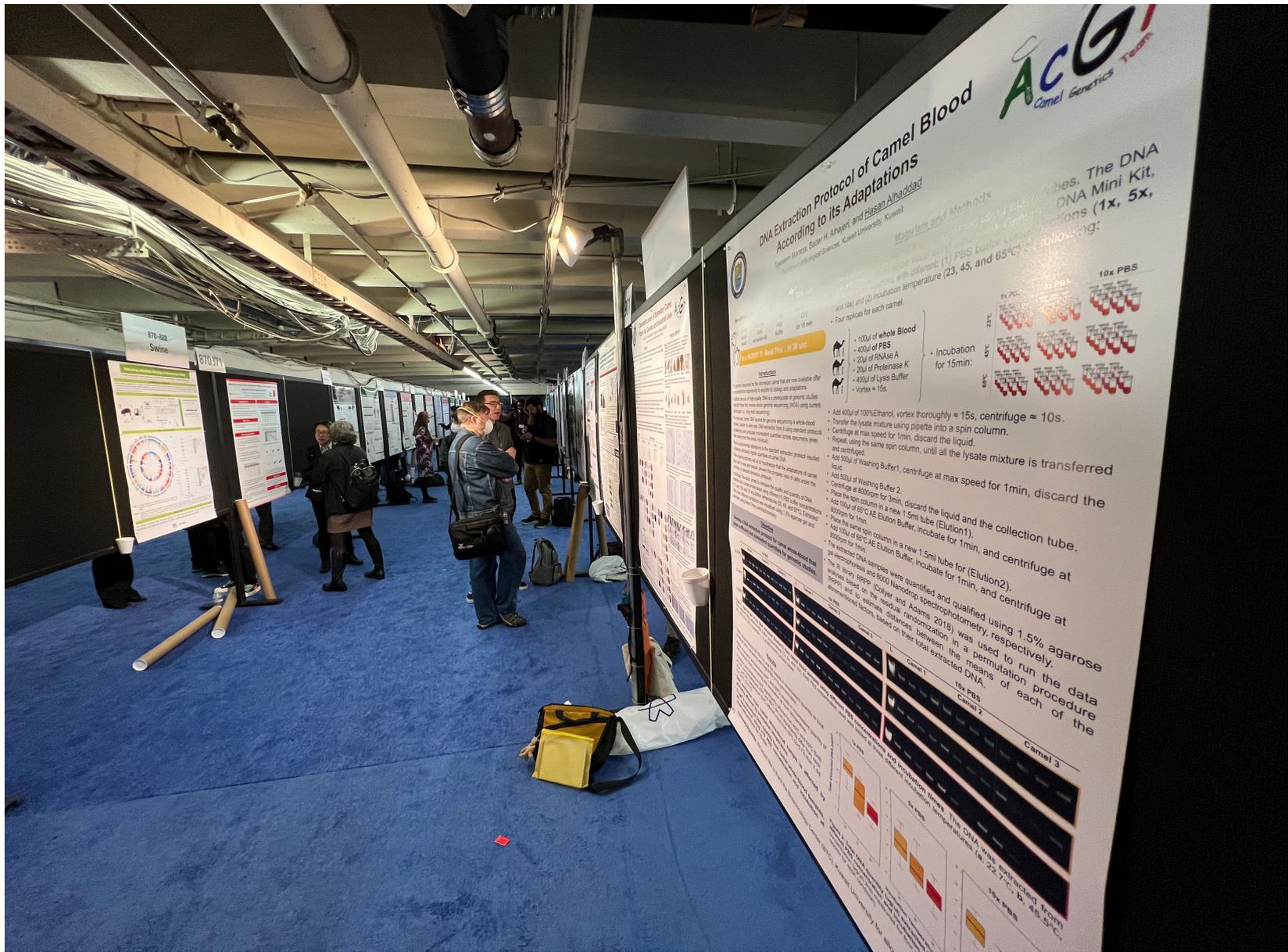
**Too much text** material is a deterrent from any short or long scientific discussion.



The general audience, people out of your field, will only glance at your poster or spend ~15-20s, if you are lucky, looking at your poster.



Colleagues within your specialized field will seek you and engage in lengthy discussion.



# DNA Extraction Protocol of Camel Blood According to its Adaptations



Youssef Marzouk, Basim H. Alhageer, and Hassan Alhaddad  
Department of Biotechnology, Kuwait University, Kuwait

**Abstract:** Camel blood has been used as a source of DNA for genetic studies. This study aims to develop a simple and efficient protocol for DNA extraction from camel blood. The protocol involves the use of a lysis buffer, followed by incubation at different temperatures (23, 45, and 65°C) for 15 minutes. The DNA is then extracted using a DNA Mini Kit, and the concentration is determined using a spectrophotometer. The results show that the protocol is effective in extracting high-quality DNA from camel blood.

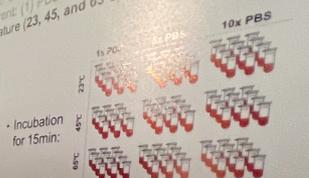
**Introduction:** Camels are one of the most resilient animals on Earth, and their blood has been used as a source of DNA for genetic studies. This study aims to develop a simple and efficient protocol for DNA extraction from camel blood. The protocol involves the use of a lysis buffer, followed by incubation at different temperatures (23, 45, and 65°C) for 15 minutes. The DNA is then extracted using a DNA Mini Kit, and the concentration is determined using a spectrophotometer. The results show that the protocol is effective in extracting high-quality DNA from camel blood.

**Materials and Methods:** The study was conducted using camel blood samples collected from three different camels. The DNA extraction protocol involved the use of a lysis buffer, followed by incubation at different temperatures (23, 45, and 65°C) for 15 minutes. The DNA is then extracted using a DNA Mini Kit, and the concentration is determined using a spectrophotometer.

**Results:** The results show that the protocol is effective in extracting high-quality DNA from camel blood. The DNA concentration was determined using a spectrophotometer, and the results show that the protocol is effective in extracting high-quality DNA from camel blood.

**Conclusion:** The study shows that the protocol is effective in extracting high-quality DNA from camel blood. The DNA concentration was determined using a spectrophotometer, and the results show that the protocol is effective in extracting high-quality DNA from camel blood.

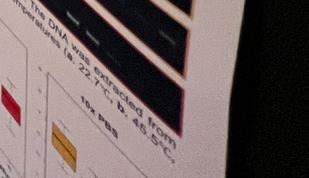
Camel blood has been used as a source of DNA for genetic studies. This study aims to develop a simple and efficient protocol for DNA extraction from camel blood. The protocol involves the use of a lysis buffer, followed by incubation at different temperatures (23, 45, and 65°C) for 15 minutes. The DNA is then extracted using a DNA Mini Kit, and the concentration is determined using a spectrophotometer. The results show that the protocol is effective in extracting high-quality DNA from camel blood.



- 100µl of whole Blood
- 400µl of PBS
- 20µl of RNase A
- 20µl of Proteinase K
- 400µl of Lysis Buffer
- Vortex = 15s.

- Add 400µl of 100% Ethanol, vortex thoroughly = 15s, centrifuge = 10s.
- Transfer the lysate mixture using pipette into a spin column.
- Centrifuge at max speed for 1min, discard the liquid.
- Repeat, using the same spin column, until all the lysate mixture is transferred and centrifuged.
- Add 500µl of Washing Buffer1, centrifuge at max speed for 1min, discard the liquid.
- Add 500µl of Washing Buffer 2, centrifuge at max speed for 1min, discard the liquid.
- Centrifuge at 8000rpm for 3min, discard the liquid and the collection tube.
- Place the spin column in a new 1.5ml tube (Elution1), 8000rpm for 1min.
- Add 100µl of 65°C AE Elution Buffer, incubate for 1min, and centrifuge at 8000rpm for 1min.
- Place the same spin column in a new 1.5ml tube for (Elution2), 8000rpm for 1min.

The extracted DNA samples were quantified and qualified using 1.5% agarose gel electrophoresis and 260/280 Nanodrop spectrophotometry, respectively. This is library RPP3P (Cotter and Adams 2016) was used to run the data analysis based on the individual randomization in a permutation procedure (RPP3P) and to estimate distances between the means of each of the determined factors, based on their total extracted DNA.



870-382 Swine

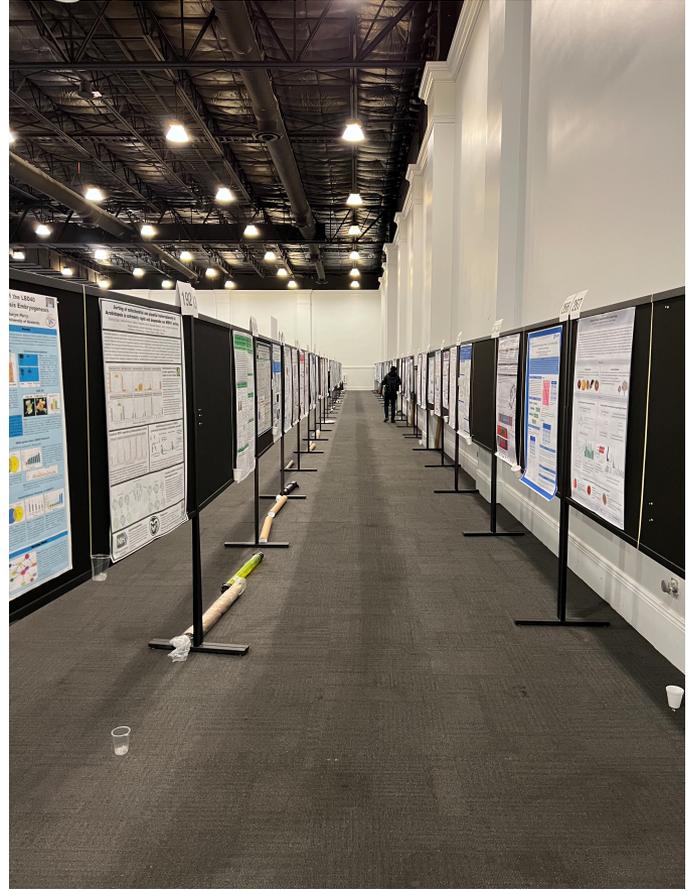
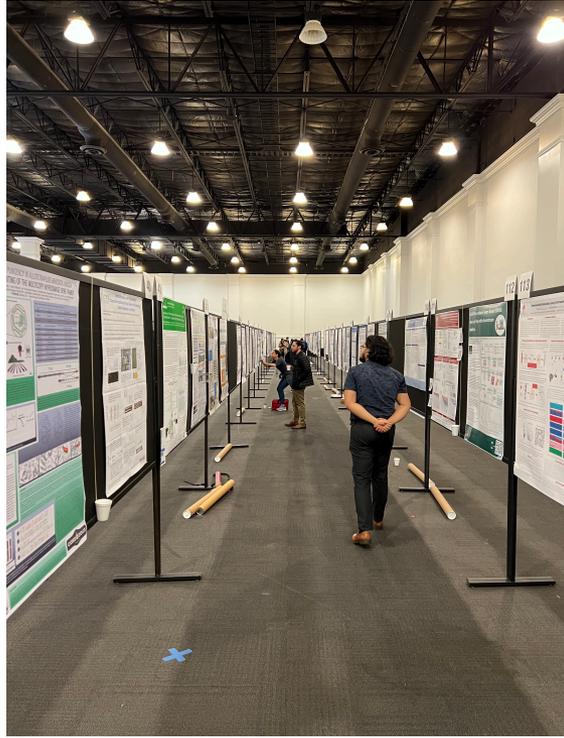
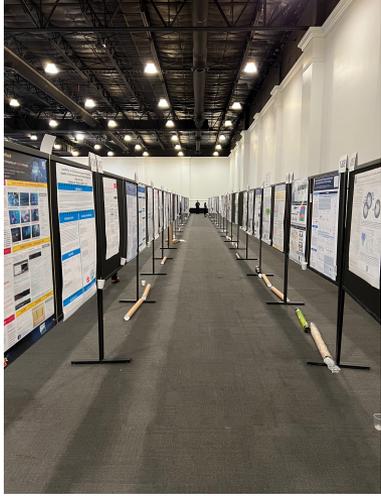


870-371





Large scientific conferences display hundreds  
if not thousands of posters.





Your goal is to catch the eyes and minds of  
as many scientists as possible.



# Design software

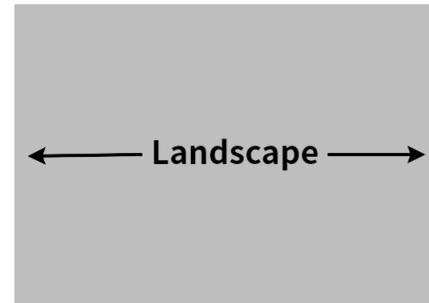
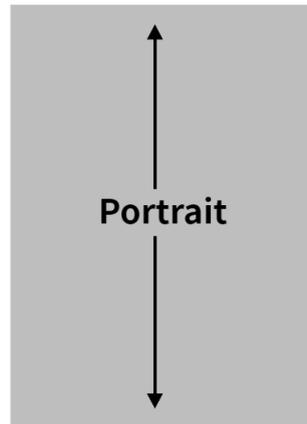


Microsoft **Powerpoint** due to its wide usage at academic institutions by faculty and students.

**IMPORTANT!**

# Layout

Both landscape and portrait layouts are used to design scientific posters.





**Portrait** layout are preferred especially in large scientific conferences since the posters' display area is limited and more portrait than landscape posters can be displayed in the same area.

**Size**

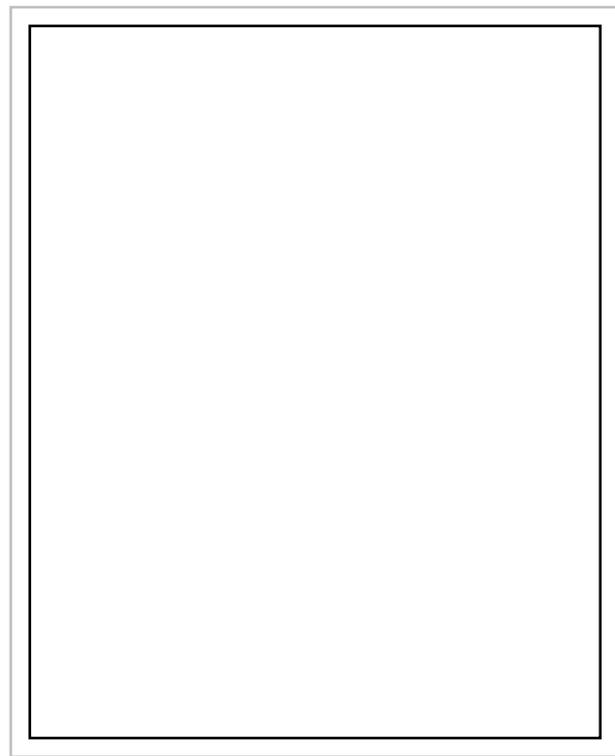


100cm height x 80 cm width  
46 inches height x 34 inches width

# Margins

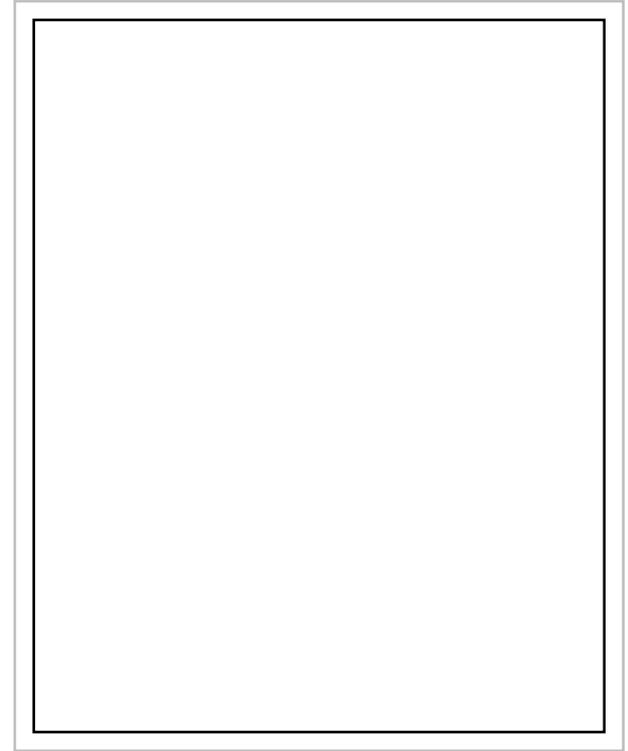


Leave  $\frac{1}{2}$  inch (1cm) margins  
from all sides of the poster.





Your poster material will be placed in 45inx33in design area (98cmx78cm).



**IMPORTANT!**

**Font type**



Use a basic font that can be easily read (ex. **Arial**).

**Arial**

**Calibri**

**Times New Roman**

**Franklin Gothic**

# Font color



You need a font color that provides the greatest color contrast with the poster background color.



Use **black** font color for most of the written content of your poster.

Use other colors for certain **words/sentences** to emphasize **importance** or a specific relationship to graphical content.

# Font size



With the exception of the authors' affiliations, figure/table legends,  
acknowledgments, and possibly the references,

**All sections should be typed with font size at  
least 28pt.**



Your poster should be readable from ~1.5m  
or 5 feet away.



You don't want to get people very close to the poster because it will reduce their interaction with you.



You don't want to get people very close to the poster because it may block other interested individuals from having visual access to your poster.

**IMPORTANT!**

**Poster background**



Avoid having a colored background or a background photo.

Consider leaving the background white.



This will concentrate the focus of the eye on the material of the poster (both writing and figures) rather than the background.



**IMPORTANT!**

**Title**



The title is the

**first most important**

component of your poster.



Use a short catchy title in a large font size  
(>60pts).



Your scientific poster is not a scientific paper to be titled with details and does not need to be easily searched and cited.



The title should be attractive to individual passing by the poster or glancing at the conference's program.



# Authors



Write the names of the authors in the order agreed upon by the research team (font size  $\leq$  36pt).

Underline the name of the presenter. The presenter may not be the first author.

This may happen when your advisor takes your work to large scientific conferences.

After each of the authors' names place a number in a superscript (e.g., Hasan<sup>1</sup>).



# Affiliations



The superscripted numbers will be the reference for the affiliations of the authors (font size ~24pt).

You can be creative with the display of the affiliations.

You do not need to give every institution/department a separate line under the author line.

You can simply separate the numbered affiliations with a comma.

# Logos



Choose good quality logos of your institution or lab.

It is preferred to choose a logo with a transparent back ground.

There is no standard location to place the logos and it all depends on the design.

I suggest that you scale the size of the logo proportionally to the height of the title lines.



# Abstract



Most scientific conferences require the submission of a ~250 words abstract for your poster, which will be printed in the conferences program or uploaded to the conference's webpage.



There is no need to include the abstract in your poster.



You can use part of your submitted abstract in the poster but not in its entirety and not in a paragraph format.

**IMPORTANT!**

# Objectives



The objectives is the

**second most important**

section of your poster.



You need to pay extra attention to the formulation of your sentences and the writing of concise and clear objectives.

The entire poster will be focused on the various ways to achieve the objectives.

Highlight this section with an attractive color and make the font bold (font size 32-36 pts).



# Hypotheses



Your hypotheses may be included in the objectives section and highlighted.

The inclusion of a hypothesis (if you have one) makes your poster easier to explain from your side and easier to read and understand by the audience.

**IMPORTANT!**

# Figures

**IMPORTANT!**

The figures are the

**third most important**

part of your poster.



Unlike your papers (written communication) or talks (oral communication), your posters are largely visual representations of your science.



Your audience will mostly look at your poster rather than read it and in many cases you will not be around to explain the content verbally.



Present your work in clear figures and diagrams that are both attractive to the eye and self-explaining to the mind. Your figures may have different forms (plots, diagrams, pictures etc.).

Generate the plots and diagrams with a transparent or white background.

This will ensure the perfect blend of your figures with the overall background of your poster.

# Tables



Avoid inserting tables in your poster.

**Table 3.1. Marker specific summary statistics of camel STR panel for the global and Mezayen datasets**

Marker	Accession No.	Repeat Motifs	Global Dataset					Mezayen Dataset				
			Geno <sup>3</sup> . Rate	Geno <sup>4</sup> . No.	Allele No.	H <sub>o</sub> <sup>5</sup>	PIC <sup>6</sup>	Geno. Rate	Geno. No.	Allele No.	H <sub>o</sub>	PIC
LCA99 <sup>1</sup>			0.99	135	36	0.89	0.88	1	66	18	0.90	0.86
CVRL01 <sup>2</sup>	AF217601	(GT) <sub>27</sub> , (GC) <sub>6</sub> , i(GT) <sub>9</sub>	0.83	102	27	0.81	0.79	1	47	23	0.70	0.70
YWLL08 <sup>2</sup>	AF217608	(CA) <sub>9</sub> , (GA) <sub>5</sub>	1.00	99	22	0.87	0.86	1	58	17	0.85	0.83
LGU75	AF237494	(GT) <sub>20</sub>	0.83	79	21	0.78	0.75	1	54	16	0.83	0.81
CMS50	AF329149	(GT) <sub>27</sub>	0.83	58	16	0.87	0.85	1	38	11	0.83	0.81
CMS121	AF329159	(TG) <sub>24</sub>	0.83	49	14	0.80	0.77	1	30	9	0.76	0.72
LCA33	AF060103	(CA) <sub>8</sub>	0.83	43	18	0.74	0.70	1	20	12	0.64	0.57
CMS9	AF329160	(GT) <sub>24</sub>	0.83	37	15	0.75	0.71	1	21	8	0.77	0.73
CVRL05	AF217605	i(GT) <sub>25</sub>	0.83	35	17	0.71	0.66	1	25	11	0.71	0.67
CMS13	AF329158	(AC) <sub>27</sub>	0.83	29	14	0.68	0.63	1	20	8	0.71	0.67
LCA66	AF091125	(CA) <sub>13</sub>	1	29	11	0.80	0.77	1	16	7	0.74	0.69
YWLL44	GU723276	(TG) <sub>18</sub>	1	24	9	0.73	0.69	1	12	5	0.63	0.58
CMS16	AF329157	(TG) <sub>34</sub>	0.83	23	12	0.60	0.53	1	16	9	0.60	0.54
LGU56	AF237492	(GT) <sub>16</sub>	0.83	20	13	0.35	0.32	1	11	6	0.45	0.41
CVRL04	AF217604	i(GT) <sub>9</sub>	0.83	13	8	0.63	0.56	1	6	3	0.63	0.55
LGU76	AF237495	(GT) <sub>11</sub>	0.83	11	7	0.65	0.58	1	8	5	0.67	0.60
VOLP32	AF305234	(TG) <sub>20</sub>	0.99	10	7	0.47	0.44	1	4	3	0.41	0.32
YWLL29 <sup>1</sup>			1	9	5	0.43	0.37	1	3	2	0.42	0.33
LCA8	AF060096	(CA) <sub>14</sub>	1	3	2	0.42	0.33	1	3	2	0.50	0.37

<sup>1</sup> Markers with unknown repeating nature. <sup>2</sup> Complex STR markers with more than one block of repeats. <sup>3</sup> Genotype rate. <sup>4</sup> Genotype number. <sup>5</sup> Observed heterozygosity. <sup>6</sup> Polymorphism Information Content.

Tables are generally detailed summary of your work or your scientific findings.

The location of such details is not your poster but your scientific papers.

**Table 3.1. Marker specific summary statistics of camel STR panel for the global and Mezayen datasets**

Marker	Accession No.	Repeat Motifs	Global Dataset					Mezayen Dataset				
			Geno <sup>3</sup> . Rate	Geno <sup>4</sup> . No.	Allele No.	H <sub>o</sub> <sup>5</sup>	PIC <sup>6</sup>	Geno. Rate	Geno. No.	Allele No.	H <sub>o</sub>	PIC
LCA99 <sup>1</sup>			0.99	135	36	0.89	0.88	1	66	18	0.90	0.86
CVRL01 <sup>2</sup>	AF217601	(GT) <sub>27</sub> , (GC) <sub>6</sub> , i(GT) <sub>9</sub>	0.83	102	27	0.81	0.79	1	47	23	0.70	0.70
YWLL08 <sup>2</sup>	AF217608	(CA) <sub>9</sub> , (GA) <sub>5</sub>	1.00	99	22	0.87	0.86	1	58	17	0.85	0.83
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LCA8	AF060096	(CA) <sub>14</sub>	1	3	2	0.42	0.33	1	3	2	0.50	0.37

<sup>1</sup> Markers with unknown repeating nature. <sup>2</sup> Complex STR markers with more than one block of repeats. <sup>3</sup> Genotype rate. <sup>4</sup> Genotype number. <sup>5</sup> Observed heterozygosity. <sup>6</sup> Polymorphism Information Content.

Tables generally occupy a large area of the poster (if displayed with a reasonable font size), which can alternatively be used for more important graphical or written content.

**IMPORTANT!**

# Conclusion



The conclusion(s) is the

**fourth most important**

section of the poster.



It should give a general summary of the findings in a concise sentence or few sentences.



The conclusions should include the general take home message of your scientific work.



# References



Your poster is not a scientific paper.  
Do you need references?

# Disclaimer

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