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Supplemental Materials

for

The Microbial Contamination of Mobile Communication Devices

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Appendix 1: Poster - Investigations Into the Microbial Contamination of Mobile Phones



Investigations into the Microbial Contamination of Mobile Phones

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Introduction



cultures taken from a toilet seat

Any is it nil by mobile'

References

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Mobile phones and other frequently handled inert items have been implicated in cross-infection in the hospital environment (see references). This practical investigation arose partly due to this scientific information, and also as a result of some media coverage.

> The Medical Microbiology module is a final year option, primarily for students taking Honours degrees in Biology and Biomedical Sciences. In the 07-08 session, 65 students enrolled. Laboratory classes comprised 4 x 4h sessions, each separated by a week (thus more than one exercise is carried out in any given session). The classes were run twice. Student: staff ratio was around 1:10.

Aim:

To demonstrate the potential of inert, frequently handled items to facilitate crosscontamination, and to encourage students to consider such issues in the hospital setting.

Objectives:

To isolate and identify common contaminants on a mobile phone, and to consider their origin, focusing particularly on presumptive staphylococci, and comparing the identity of those isolated from the phone with those isolated from the skin.

Methods

The schedule was provided to students in advance of the class. Students record observations on the gapped schedule, and subsequently download new sheets from WebCT to produce the final report. The report was divided into two sections: the first required students to outline observations and explain underlying principles of microbiology via a 'gapped' question sheet; the second asked students to assess the source (skin flora?), and significance of contamination of inert objects in terms of cross-infection, particularly within the hospital setting.

A shortened version of the schedule follows:

WEEK 1: ISOLATION

Work individually.

- Moisten a sterile swab in saline and swab your skin (eg hands, 1. Take digital pictures of the mobile phone plates which you nose, ears, lips). Roll the swab firmly over the surface. Use the swab to inoculate a tryptone soy agar plate, using the isolation plate technique to ensure separation of colonies.
- Draw a diagram to show how you inoculated the plate. Firmly press your mobile phone (if you have one) onto a large agar plate OR if you have a flip phone, use a moistened swab to sample the suface (particularly those that come into contact with mouth, ears, fingers). Inoculate the plate with the swab to ensure maximum yield of colonies. Draw a diagram to show how you inoculated the plate.
- Use the medicated wipes provided to clean your phone. Then resample (either by pressing onto agar, or by swabbing, as before). Wipe the phone again.

WEEK 2: PURIFICATION

Incubate plates 37° 24 h

- Take digital pictures of the mobile phone plates which you inoculated last week.
- REPEAT THE PHONE SAMPLING ie press or swab, before and after cleaning. (Wipe clean again after second sampling)
- Looking at last week's plates, describe any differences between the microflora of the two sites (phone pre-clean, and skin) and explain why they occur.
- It is likely that some of these colonies will be staphylococci. Why might this be? Have you got any colonies which are 'presumptive staph' on both skin and phone?
- What are the typical colony morphologies of staphylococci (text book)?
- Subculture one colony of each of your presumptive staphylococci onto mannitol salt agar and nutrient agar to obtain pure **For your report:** cultures on selective and non selective media for identification ('isolation plates'). What is in mannitol salt agar, and why is it used?
- Why do you usually need a pure culture on a non-selective medium (as opposed to a selective medium) for identification?
- What other colonies were present on your mobile phone and skin swabs? Do Gram stains, and suggest identification and sources of these organisms.

WEEK 3: IDENTIFICATION

- inoculated last week. REPEAT THE PHONE SAMPLING – ie press or swab, before and
- after cleaning. Wipe clean after sampling. 2. Look at the isolation plates you set up last week. Check the cultures are pure. Do the colonies look like those you subcultured? Use the non-selective medium from now on.
- You must have pure cultures for the identification test. To confirm Staphylococcus genus, perform Gram stain and catalase test.
- For the following tests you must have a positive control (S.aureus). Lab cultures are provided.
- To identify species, perform
- -- latex aggulation test (for coagulase and protein A confirmatory for S.aureus)
- -- API staph for coagulase negative staphylococci (CNS) only. Construct a table summarising the properties of your isolates,
- and state their identities. Paste in any API printouts (results week 4)
- ANTIBIOTIC SENSITIVITIES Set up a plate to show antibiotic
- sensitivities of two of your isolates (eg one from skin and one from phone, with similar colony morphologies).

WEEK 4: FINAL WORK

Take digital pictures of the mobile phone plates which you inoculated last week.

Identify any isolates via API system. Measure zones of inhibition for antibiotic sensitivities.

Insert photos from the three cleaning protocols (agar plates before and after cleaning).

Comment on the experiment and the results you obtained. Can you comment on mobile phone contamination (extent, origin, type of microorganisms, ease of cleaning, importance of cleaning, buildup of contamination etc.)? Do you see any hygiene or cross-contamination issues in this context? Were there differences between phones/people? Compare drugs resistance and other results of your isolates and discuss implications. Are you concerned? Are you a carrier?

What other inanimate objects might pose comparable crosscontamination issues? Look for references illustrating similar work.

Discuss in the light of lectures and literature. State references.

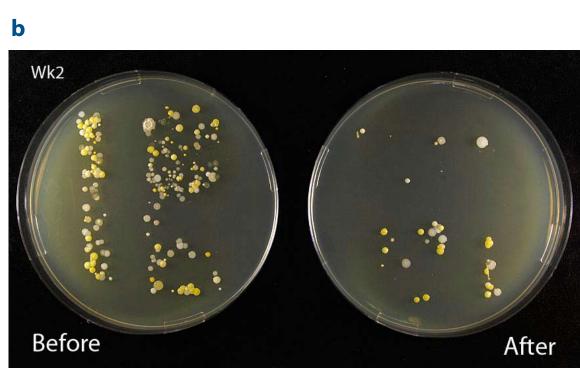
Results

Antimicrobial wipes reduced contamination, particularly on smooth areas of the phone. Not all wipes were antibacterial. [Fig 2].

Some students were able to isolate similar species of staphylococci from phone and skin. Most isolated micrococci from their phones. Bacillus spp. were also isolated. None isolated Staphylococcus aureus

Figure





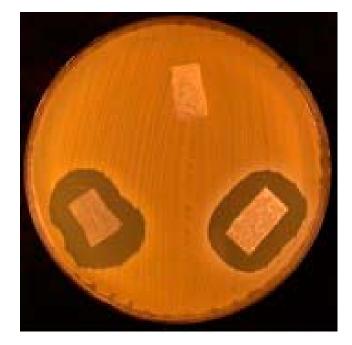


Of the 63 stude	nts who sub	mitted work	, marks were	as follows:	
Mark (%)	0 - 20	21 - 40	41 - 60	61 - 80	81 - 100
No. students	2	10	24	19	8
Low marks were awarded due to non-completion of the gapped questions and an absence of an exploratory/explanatory text.					

All students were able to demonstrate contamination of phones, and recognised the transient and nature of the contamination (variation from week to week; comparison with other students). [Fig 1 a-c].



Figure 2



Evaluation

The exercise was evaluated by written questionnaire, to which 38 responses were provided.

Responses to Question

(What did you think of the mobile phone practical overall?)

Positive	Negative
Good	Long and unorg
It was interesting and simple	Not very good
Practical was good- learned how to identify colonies	
I really enjoyed the practical it was very interesting	
It was a good practical	In between
lt was good, l enjoyed it, it was fun	Ok, too long
Interesting practical, fun, visual bacteria	Ok
Very good	
Exciting, enjoyed every bit of the practical	
It was a good idea, esp. since everybody has a mobile now	
Was good and informative	
It was well presented and easy to understand, the lecturers managed to convey their enthusiasm for the subject matter well	
Very interesting	
Good but very lengthy with lab report, was very in- teresting though. Practicals should have been worth more though	
Was good and interesting to do but was also quite all over the place with several parts not clearly outlined	
Quite interesting to actually see the flora present on an inanimate object instead of just being told they were there	
It was interesting to see the range of organisms that are carried on the skin and those which are picked up from the environment	
Informative	
Confusing, but overall interesting	
l enjoyed it but the timing was rubbish as we had our dissertation deadline then too	
Interesting, possibly best practical of my third year	
It was interesting though hard to get results	
Very interesting & enjoyable	
It was interesting, possibly not as challenging as the other practicals we were set. Instead of a write-up could do a worksheet?	
Very good as it linked in with what we were studying in the lectures and helped in learning	
Very positive	
Interesting, made it more fun to do, learnt more that way	
Very good- alerting, informative and interesting	

Responses to Question 2

(What was the best aspect?)

Practical based	
Just seeing the sheer amount of different bacterial species that were phone	on y
Its hands on and really does show that microbes are part of everyday	y life
Seeing what grew on the plate after week 1 (was pretty disgusting)	
Finding how much bacteria was on your phone	
The phone itself	
Seeing what was on own item	
Visualising the organisms	
Realising just how much bacteria is present on the mobile	
Seeing the pictures	
Visualisation and isolation of organisms	
Learning new diagnostic tests	
Detecting the normal flora	
Culturing imprint of mobile phone to see what is found on it	
Getting to identify organisms on phone and compare with skin	
Seeing the results	
Seeing if I carried any potentially pathogenic organisms	
Plates stamping	
Eye-opener, made you question possible contaminants and what ma Personal- using own mobile phone gave a fun, personal approach	ay be
Practical	
Seeing what organisms are found on a phone	
Plating the phones	
Discovering how contaminated phones can be, as well as the whole taminants possible	varie
Realising how much my phone was contaminated!	
API strips	
Everything was really good, especially looking at the amount of gerr relationship between phone and skin	ns pi
All the staff in the unit were much more helpful than any other unit	
It gave a good insight to microbial skin populations and their differe	nt ty
Different, learnt stuff that applied to you- as you used your own mob	oile

Different, learnt stuff that applied to you- as you used your own mobile Seeing what we were carrying, and how the phone became contaminated over the





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	More stringer and comparin if antibacteria
	More help ide
	Should be wo

esponses to Qu	lestion 3
What parts could	d have been improved?)

Mana time situan. Thus a prostical to prote success with difficult. No shad to be shown
More time given. Three practical together was quite difficult. Needed to be slower/ more spaced out so that could understand what was going on! VERY RUSHED
It got a bit hectic some weeks doing 3 pracs all at the same time
I think the length of time but that can not be changed due to obvious incubation times
Not sure really I thought it was pretty well organised
All parts
Mine was really boring with a few dots
The lab reports are only worth 10% each and I think there was a lot of work needed for each report
Too many practical's on the go at once
More stringent controls on which wipes used. Perhaps setting up different groups and comparing the effectiveness of certain types of wipes. For example to find out if antibacterial wipes are worth paying extra for
More help identifying bacteria
Should be worth more % for detail required
The organisation and layout of work and experiments
Isolation/culturing of pure isolates proved tricky
I don't think it needed to be done every week
Write-up too difficult
Separation from other experiments/practicals been run, confusing to know why w were doing what we did
Not have such a huge write-up when other deadlines (dissertation) are due in
The queuing for photos
This year the practical was hectic, having 3 practicals on the go each week made it difficult
Time, number of students per prac decreased
I thought there was a bit too much going on in the practical at one time
A group demonstration at the end of the practical to see what everyone found and the changes
Too many things all added together- had to do other practicals too, could spread it out more not mix up different practicals in one day-confusing
Storage of the plates-some became lost. Also the last session was INCREDIBLY rushed- very stressful!!

Responses to Question 4

(Do you think the mobile phone practical would make a good exercise in future medical microbiology units?)

Positive	Negative
Yes	No
Yes, it was extremely interesting and a bit more fun	Definitely not
Definitely	
Yes. I learn things better if I enjoy them I'm sure that's true to others too	
Yes, it was interesting	
Yes I do	In between
Yes, because it demonstrates the fluidity of what's on dif- ferent peoples phones	Possibly if results could be guaranteed
Yes. Its interesting to see the range of organisms we come into contact with everyday	
Yes- improved write-up theory	
Definitely! Once over the confusion, was great learning tool	
Possibly if results could be guaranteed	
Yes as I really enjoyed it	

Conclusions

As with all new lab classes, several improvements can be made to its efficient and effective operation. However, students clearly enjoyed the exercise, and it was apparent that learning had taken place.

The class will be run in 08-09, with an alteration in assessment emphasis (inclass assessment only of other concurrent experiments, with the mobile phone report as the key laboratory assessment)