



Device Choices for OPAT – stability data and patient safety

Tim Hills

Structure

Background/Nottingham OPAT approach

Administration devices

Drug stability

BSAC project

Nottingham OPAT Approach

- Clinical service started October 2011
 - Patient self-admin
 - Small infusion centre
- March 2012
 - Pre-filled devices via private healthcare company
 - HCW administered via private healthcare company

Options for Self-administration

- Syringes, diluents and vials
- Pre-compounded syringes
- Bags, flow restrictors/pumps
- Elastomeric devices
- 24hr Syringe pumps
- N.B. Pro/Cons Nottingham opinion

Infusion Bags and Pumps

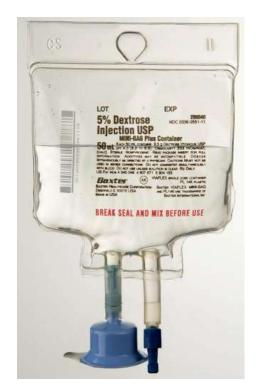


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- Flow restrictors
 - Often poor tolerance values for accurate administration.
 - OK for standard infusions (e.g. cephalosporins, penicillins, carbapenems)
 - Not for vancomycin/ drugs where rate important.
- Electronic Pumps and giving set.
 - Can be harder to learn
 - More accurate

Closed-admixture bags

- Simplifies number of steps.
- Still requires dexterity for attaching giving-set.
- Flow-restrictor/pump still needed.
- Option if stability short.



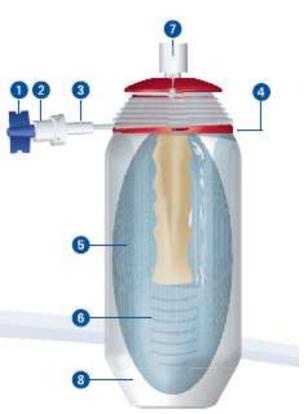


Elastomeric Devices

- HomepumpEasypump©/Eclipse©
- Intermate ©/Infusor©
- Reasonable options of all OPAT drugs (inc. vancomycin).
- 24hr infusion option available



Large Volume Infusor Small Volume Infusor



Factors Affecting Elastomeric Flow Rates

- Built-in flow restrictor- main variable as per manufacturer
- Viscosity Temperature 10°C change can →
 20-30% viscosity change check calibration temps.
 - 100ml 6 hours out of fridge before use (Home pump eclipse[©])
 - 6 hours Intermate[©] large volume
 - 4 hours Intermate[©] small volume

Factors Affecting Elastomeric Flow Rates

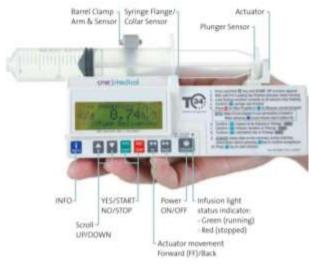
- Pressure gradient
 - Underfill/overfill

Potential overall fluctuation of +/-40%

Patient education on correct use essential.

Syringe Drivers

- Potential option for 24 hour infusion via syringe.
- Generally harder to learn compared with elastomeric.
- Consider for "infusion-centre" model.



NUH service (Oct12-Apr13)

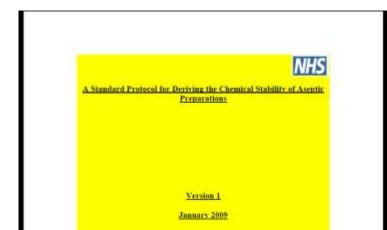
Method of administration	% of total episodes (6 months)
Taught – Self/Carer administered	76%
Taught using prefilled	10%
HCW delivered in patient home	4%
Ward based infusion centre	10%

Obtaining prefilled devices

- Local pharmacy sterile production department
 - Under supervision of a pharmacist "special" (max 7 days)
 - Manufacturing licence (longer expiry but batch manufacturer and QC)
 - Penicillins
- Other NHS Trust production department
- Private/Homecare provider

"Appropriate" stability data

- "Special" licenced
 - Dependent on opinion of specialist production pharmacist.
- "Batch production"
 - In agreement with local QC department (National QC standards 2009).



BSAC OPAT Drug Stability in Devices Project - Aims

Remove barriers to local production through:

Review stability literature

Identify gap
 commission stability studies

BSAC OPAT Drug Stability in Devices Project - Tasks

- 1. Determine which drugs/devices centres would be keen to use.
- 2. Review the literature around stability.
- 3. Undertake stability testing to national standards where gaps in the literature exist and publish in the public domain.



https://www.surveymonkey.com/s/opatdrugstability





Drug Stability & Testing in the OPAT Setting		
Drugs & Devices - Would Consider Using		
Please indicate which drugs & devices you would consider using in	n the OPAT setting.	
8. Which drugs would you consider using in a prefilled syrin	ge (produced by an SPU) in the OPAT setting (if it wa	s available)?
	Would consider	Would not consider
Aztreonam (2g)	0	O
Benzylpenicillin (12g over 24hrs)	0	0
Benzylpenicillin (14.4g over 24 hrs)	0	0
Ceftazidime (1g)	0	0
Ceftriaxone (1g)	0	0
Colistin (2mu)	0	0
Meropenem (1g)	0	0
Meropenem (3g over 24 hrs)	0	0
Meropenem (6g over 24 hrs)	0	0
Tazocin (4.5g)	0	0
Tazocin (13 5g over 24hre)		0

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