

Patient trial number: 14-



#### UKALL14 Pregnancy Report

### NOT REQUIRED FOR 'REGISTRATION ONLY' SUB-STUDY PATIENTS (14-3-XXX)

Please fax this form to the UKALL14 Co-ordinator at the CR UK & UCL Cancer Trials Centre on 020 7679 9861 within 24 hours of becoming aware of the pregnancy.

Please complete all sections with details of any pregnancy occurring from the time of informed consent until last follow-up visit.

| Trial details  |  |  |                 |  |  |  |  |  |  |
|--|--|--|-----------------|--|--|--|--|--|--|
| Trial title  | A randomised trial for adults with newly diagnosed acute lymphoblastic leukaemia |  |                 |  |  |  |  |  |  |
| Trial acronym  | UKALL14  | EudraCT number   | 2009-012717-22  |  |  |  |  |  |  |
|  |  |  |                 |  |  |  |  |  |  |
| Patient details (Any information regarding female partners of trial patients should be entered in Other Pregnancy Information section) |  |  |                 |  |  |  |  |  |  |
| Patient initials   |  | Patient trial number Randomised patients only 14-1-XXX or 14-2-XXX Pregnancy data not collected for 'Registration only' patients (14-3-XXX)                                      |                 |  |  |  |  |  |  |
| Report relates to:   | Trial Patient Patient's Partner  | Patient date of birth  |                 |  |  |  |  |  |  |
| Hospital   |  | Treating Clinician   |                 |  |  |  |  |  |  |
| B Randomisation:   | 0=N/A 1=B1 2=B2 <b>T Randomisation</b> : 0=N/A 1=T1                              | 2=T2 P Randomisation:  | 0=N/A 1=P1 2=P2 |  |  |  |  |  |  |
| Type of report   | Initial d d m m m y y  | For all follow up reports, please:  Initial & date all changes throughout the report  Follow up  Fax to UCL CTC within 24 hours of becoming aware of significant new information |                 |  |  |  |  |  |  |
| Complete for initial reports only:   | Date site notified of pregnancy:  d d m m m y y                                  | If reported to UCL CTC more than 24 hours after becoming aware of pregnancy, provide reason:   |                 |  |  |  |  |  |  |





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|--|-------------------------------|-----------|-------|-------------|---------------------|--------------|----------------------|---------------------------|---------------------------|--|
| Trial treatment  |                               |           |       |             |                     |              |                      |                           |                           |  |
| Drug Name  | Brand                         | Dose      | Unit  | Frequenc    | Is this full dose?  | Route        | Start date           | Ongoing?                  | End date                  |  |
| Pegylated asparaginase   | Oncaspar                      |           |       |             |                     | N            |                      | $\square_{Y} \square_{N}$ |                           |  |
| Rituximab  | Mabthera                      |           |       |             |                     | N            |                      | Y N                       |                           |  |
| Nelarabine   | Atriance                      |           |       |             |                     | N            |                      | $\square_{Y} \square_{N}$ |                           |  |
| Palifermin   | Kepivance                     |           |       |             |                     | N            |                      | Y N                       |                           |  |
| Most recent phase of protocol treatment (1=Phase 1 induction, 2= Phase 2 Induction, 3= Intensification, 4= Consolidation, 5= Maintenance, 6= Myeloablative transplant, 7= Non-Myeloablative transplant)  Date last treatment given before pregnancy confirmation:  Confirmation:  Last trial drug given before pregnancy confirmation: |                               |           |       |             |                     |              |                      |                           |                           |  |
|  |                               |           | (Only | includa dru | as aiven during the | 20 day nario | d prior to programou |                           |                           |  |
| Concomitant medications?  (Only include drugs given during the 30 day period prior to pregnancy confirmation. Continue on separate sheet if necessary)  Continued on a separate sheet:   |                               |           |       |             |                     |              |                      |                           |                           |  |
| Drug Name  | Brand                         | Indicatio | on    | Dose U      | nits Frequenc       | y Route      | Start date           | Ongoing?                  | End date                  |  |
|  |                               |           |       |             |                     |              |                      | Y N                       |                           |  |
|  |                               |           |       |             |                     |              |                      | $\square_{Y} \square_{N}$ |                           |  |
|  |                               |           |       |             |                     |              |                      | Y N                       |                           |  |
|  |                               |           |       |             |                     |              |                      | Y N                       |                           |  |
|  |                               |           |       |             |                     |              |                      | Y N                       |                           |  |
|  |                               |           |       |             |                     |              |                      | Y                         |                           |  |
|  |                               |           |       |             |                     |              |                      | Y N                       |                           |  |
|  |                               |           |       |             |                     |              |                      | $\square_{Y}\square_{N}$  |                           |  |





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| Fatient trial number:                                       |   |                         |                          |  |                |    |                             |   |  |  |
|---|---|-------------------------|--------------------------|--|----------------|----|-----------------------------|---|--|--|
| Pregnancy Information                                       |   |                         |                          |  |                |    |                             |   |  |  |
| Start date of last menses                                   | Date pregnancy confirmed                |                         |                          | Method of diagnosis  |                |    | ticipated date of childbirt | Mother consented for pregnancy monitoring |  |  |
| d d m m m y y   | d d m m m y y                           |                         |                          |  |                |    | d d m m m y y               |   |  |  |
| If consented for pregnancy monitoring:                      |   | consent signed          | PIS version used: .      |  |                |    | Consent form version:       |   |  |  |
| * If mother has not yet consented for pregnancy monitoring: | nsented for pregnancy     Will be conse |                         | c visit Other (specify): |  |                |    |                             |   |  |  |
| Pregnancy Outcome   |   |                         |                          |  |                |    |                             |   |  |  |
| Not known at this date                                      |   | Still birth             |                          | Induced abortion   |                |    | Spontaneous abortion        |   |  |  |
| Neonatal death  |   | Uneventful (norma baby) | l/healthy                | Birth defects (provide details in Other Pregnancy Information section below) |                |    |                             |   |  |  |
| Date of Above Outcome:                                      |   | d d m m m y             | у                        |  |                |    |                             |   |  |  |
| Date of delivery  | Gestation (weeks)                       | Mode of Delivery        |                          | Sex  | Weight<br>(kg) | An | tenatal Problems            | Postnatal Problems                        |  |  |
|   |   |                         | М                        | ale Female   |                |    |                             |   |  |  |





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| Patient trial number: 14  |                      |                  |         |             |  |                 |  |                    |                |                    |         |
|---|----------------------|------------------|---------|-------------|--|-----------------|--|--------------------|----------------|--------------------|---------|
| Other Pregnancy Information (concurrent conditions, medical history, complications during birth, birth defects etc) |                      |                  |         |             |  |                 |  |                    |                |                    |         |
|   |                      |                  |         |             |  |                 |  |                    |                |                    |         |
|   |                      |                  |         |             |  |                 |  |                    |                |                    |         |
|   |                      |                  |         |             |  |                 |  |                    |                |                    |         |
|   |                      |                  |         |             |  |                 |  |                    |                |                    |         |
|   |                      |                  |         |             |  |                 |  |                    |                |                    |         |
|   |                      |                  |         |             |  |                 |  |                    |                |                    |         |
|   |                      |                  |         |             |  |                 |  |                    |                |                    |         |
| Past Pregnancy History  |                      |                  |         |             |  |                 |  |                    |                |                    |         |
| Date of delivery  | Gestation<br>(weeks) | Mode of Delivery |         | Sex         |  | Weight<br>(kg)  |  | Antenatal Problems |                | Postnatal Problems |         |
|   |                      |                  |         | Male        | Female                                     |                 |  |                    |                |                    |         |
|   |                      |                  |         | Male        | Female                                     |                 |  |                    |                |                    |         |
|   |                      |                  |         | Male        | Female                                     |                 |  |                    |                |                    |         |
| <b>Signature:</b> PI or other participating clinicians only   |                      |                  |         | Print name: |  |                 |  |                    | Date of report | d d m              | m m y y |
| Office use only   |                      |                  |         |             |  |                 |  |                    |                |                    |         |
| Trial Reference: 14 CTC Reference: PREG   |                      |                  |         |             |  |                 |  |                    |                |                    |         |
| *Reported to MHRA:  | rted to Main REC     | d d m            | m m y y |             |  | Entered on data |  | m m m y y          |                |                    |         |
| Form checked by (signature)   | Date d m m m y y     |                  |         |             | Checked by clinical reviewer d d m m m y y |                 |  |                    |                |                    |         |