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Dear Professor Jureidini

Thank you for your letter to Sir Andrew's office which was passed on to me.

I have checked on the status of your enquiry about access to anonymised patient level data from paroxetine Study 329.

Information for this study and other studies of paroxetine in paediatric and adolescent patients is available on our external website, <a href="www.gsk.com">www.gsk.com</a>. The following information can be accessed directly from this link: - (<a href="http://www.gsk.com/media/resource-centre/paroxetine/paroxetine-paediatric-and-adolescent-patients.html#reports">http://www.gsk.com/media/resource-centre/paroxetine/paroxetine-paediatric-and-adolescent-patients.html#reports</a>):

- Clinical Study Report(s)
- Protocol and blank Case Report Form (Appendix A)
- Patient data listings (Appendices B to H). This contains the information directly from the case report forms.

In addition, I can confirm that we are able to provide access to the anonymised raw datasets from this study in an electronic database as described in our request site. In order for you to gain access to this we would ask that you complete and submit a formal research proposal via the request site which will be reviewed by the Independent Review Panel. A decision from the panel can be expected within 30 days (usually 5-10 days) of receipt of your research proposal. In addition we will require a signed Data Sharing Agreement. The Agreement template is also available via the request site. Given the age of this study we anticipate that anonymisation of the data may take 4-6 weeks after we receive a research proposal that is approved by the Independent Review Panel.

I hope that this is satisfactory and if you have additional questions please do not hesitate to contact me directly.

Yours sincerely

James Shannon Chief Medical Officer

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